

DESCRIPTION

NOVEL BENZOFURAN DERIVATIVE, MEDICINAL COMPOSITION CONTAINING
THE SAME, AND USES OF THESE

TECHNICAL FIELD

5 The present invention relates to novel benzofuran derivatives, which exhibit adenosine A_{2A} receptor antagonistic activities, pharmaceutical compositions containing the same, and their uses.

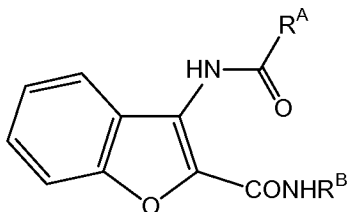
BACKGROUND ART

10 Adenosine is an endogenous purine nucleoside, and has a wide variety of regulatory functions and physiological effects. Four distinct adenosine receptor subtypes are known as A₁, A_{2A}, A_{2B} and A₃ receptors. Most of the actions of adenosine are mediated by the interaction with its membrane receptors: A₁, A_{2A}, A_{2B} and
15 A₃ receptors which belong to G-protein coupled receptor families.

Distribution and function of adenosine A_{2A} receptors in the central nervous system are well known. It has been reported that adenosine A_{2A} receptors are involved in the regulation of cholinergic, GABA-nergic and glutamate neurons. Moreover, it
20 has been reported that adenosine A_{2A} receptors are associated functionally with dopamine D₂ receptors, and blockade of adenosine A_{2A} receptors increases the binding affinity of dopamine to dopamine D₂ receptors (see nonpatent literature 1). Parkinson's disease is well known as a disorder relevant to
25 impairment of dopamine neuron. Parkinson's disease is a progressive neurodegenerative disease, which usually affects elderly patients and is characterized by impairment in

coordinated motor function such as rest tremor, rigidity, akinesia, postural instability and the like. It is thought that Parkinson's disease results from deficiency of dopamine in the striatum, which is caused by degeneration of dopamine neuron
5 in the substantia nigra. Adenosine A_{2A} receptors are abundantly present in the striatum which is playing a crucial role in the regulation of coordinated motor function. As discussed above, there is an opposing relationship between adenosine A_{2A} receptors and dopamine D2 receptors. Accordingly, it is expected that
10 agents selectively antagonizing adenosine A_{2A} receptors are useful for treating motor function disorders such as Parkinson's disease, Huntington's disease, Wilson's disease and the like (see nonpatent literatures 2 to 4). It has been reported that blockade of adenosine A_{2A} receptors results in antidepressing,
15 anxiolytic or neuroprotective actions. Accordingly antagonists of adenosine A_{2A} receptors are expected to be useful for treating depression, anxiety, cognitive function disorders such as Alzheimer disease and the like (see nonpatent literatures 5 and 6, patent literature 1). It has also been reported that blockade
20 of adenosine A_{2A} receptors ameliorates disorders after cerebral ischemia and reduces the volume of cerebral infarction. Accordingly, antagonists of adenosine A_{2A} receptors are expected to be useful for treating cerebral ischemia disorders such as cerebral apoplexy, brain lesion after cerebral vasospasm and
25 the like (see nonpatent literature 7). Antagonists of adenosine A_{2A} receptors are also expected to be useful for treating restless legs syndrome (see patent literature 2).

Sangapure S.S. and Agasimundin Y.S. et al disclose benzofuran derivatives of the following general formula:



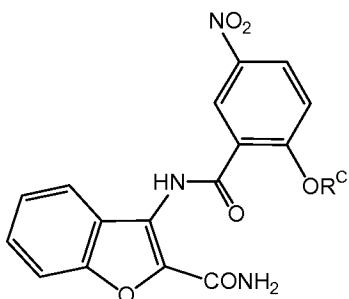
wherein R^A is a phenyl group, unsubstituted or substituted with halogen, lower alkyl, lower alkoxy or carboxy, a benzyl group or a lower alkyl group; R^B is a hydrogen atom or a lower alkyl group, as an intermediate for preparing benzofuro[3,2-d]pyrimidine derivatives (see nonpatent literatures 8 to 11). However, with regard to biological activities of the benzofuran derivatives, there is no description at all.

Basavaraj P. et al disclose 1-acetylamino naphtho[2,1-b]furan-2-carboxamide and 1-benzoylamino naphtho[2,1-b]furan-2-carboxamide as an intermediate for preparing 4-oxonaphtho[2,1-b]furo[3,2-d]pyrimidine derivatives which are useful as an antimicrobial agent, anthelmintic agent or anti-inflammatory agent (see nonpatent literature 12). However, with regard to biological activities of the naphtho[2,1-b]furan derivatives, there is no description at all.

3-Benzoylamino-5-chlorobenzofuran-2-carboxamide, 5-chloro-3-[2-(3,4-diethoxyphenyl)acetylamino]benzofuran-2-carboxamide, 5-bromo-3-[2-(3,4-diethoxyphenyl)acetylamino]benzofuran-2-carboxamide, 5-chloro-3-(2-chloroacetylamino)benzofuran-2-carboxamide, and 3-acetylamino-5-chloro-

benzofuran-2-carboxamide are registered in Chemical Abstract (see nonpatent literatures 13 to 17). However, with regard to biological activities of these compounds, there is no description at all.

5 Oota T. et al disclose benzofuran derivatives represented by the following general formula:



wherein R^C is a lower alkyl group, as an intermediate for preparing benzofuro[3,2-d]pyrimidine-4-one derivatives which are useful as a phosphodiesterase inhibitor specific to cyclic GMP (see
10 patent literature 3). However, with regard to biological activities of the benzofuran derivatives, there is no description at all.

Nonpatent literature:

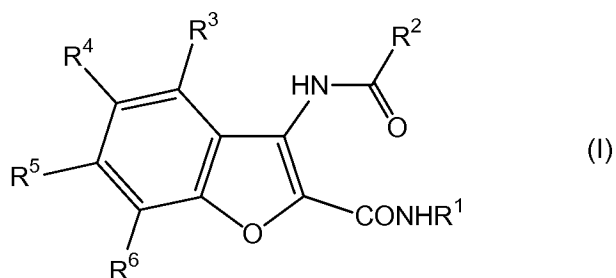
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15 p.7238-7241
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 13. "Chemical Abstract", Registry Number 340017-67-6
 14. "Chemical Abstract", Registry Number 663931-40-6
 15. "Chemical Abstract", Registry Number 633288-47-8
 16. "Chemical Abstract", Registry Number 397881-00-4
 17. "Chemical Abstract", Registry Number 332375-01-6
- Patent literature:
1. International Publication No. WO2004/108137 pamphlet
 2. International Publication No. WO2004/019949 pamphlet
 3. Japanese Patent Laid Open No. 1995/267961

DISCLOSURE OF THE INVENTION

The present inventors have intensively investigated a novel compound having adenosine A_{2A} receptor antagonistic activities, and found surprisingly that benzofuran derivatives represented by general formula (I) show a potent adenosine A_{2A} receptor antagonistic activity, and are useful for treating or preventing a disease mediated by adenosine A_{2A} receptors. Based on these findings, the present invention has been accomplished.

The present invention therefore provides a compound represented by general formula (I):



a prodrug thereof, or a pharmaceutically acceptable salt thereof, wherein

R¹ is a hydrogen atom or a lower alkyl group;

R² is:

- a) a lower alkyl group,
- b) a halo-lower alkyl group,
- c) a hydroxy-lower alkyl group,
- d) a cycloalkyl group,
- e) an aryl-cycloalkyl group,
- f) a heterocycloalkyl group,
- g) an aryl group, unsubstituted or substituted with 1 to 5 substituents selected from the group consisting of X¹, X², X³,

X^4 and X^5 ,

h) an aralkyl group, wherein the ring of the aralkyl group is unsubstituted or substituted with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 ,

5 i) an aryl-alkenyl group, wherein the ring of the aryl-alkenyl group is unsubstituted or substituted with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 ,

j) a lower alkyl group substituted with a group selected from a lower alkoxy group or a lower acyloxy group,

10 k) an aryloxy-lower alkyl group, wherein the ring of the aryloxy-lower alkyl group is unsubstituted or substituted with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 ,

15 l) an aralkyloxy-lower alkyl group, wherein the ring of the aralkyloxy-lower alkyl group is unsubstituted or substituted with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 ,

20 m) an arylsulfanyl-lower alkyl group, wherein the ring of the arylsulfanyl-lower alkyl group is unsubstituted or substituted with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 ,

n) a heteroaryl group, unsubstituted or substituted with 1 to 3 substituents selected from the group consisting of X^6 , X^7 and X^8 , or

25 o) a heteroaryl-lower alkyl group, wherein the ring of the heteroaryl-lower alkyl group is unsubstituted or substituted with 1 to 3 substituents selected from the group consisting of

X^6 , X^7 and X^8 ;

X^1 , X^2 , X^3 , X^4 and X^5 are each independently:

- a) a halogen atom,
- b) a lower alkyl group,
- 5 c) a halo-lower alkyl group,
- d) a cycloalkyl group,
- e) a lower alkoxy group,
- f) a halo-lower alkoxy group,
- g) a cycloalkyloxy group,
- 10 h) a heterocycloalkyloxy group,
- i) a lower alkoxy-lower alkoxy group,
- j) a hydroxy-lower alkyl group,
- k) a hydroxyl group,
- l) a carboxy group,
- 15 m) a lower alkoxycarbonyl group,
- n) an aralkyloxycarbonyl group,
- o) a lower acyl group,
- p) a cyano group,
- q) $-A^1-NR^{20}R^{21}$,
- 20 r) $-A^2-SR^{22}$,
- s) $-SO_2NR^{23}R^{24}$,
- t) a phenyl group, unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group, a lower alkoxy group, a halo-lower alkoxy group and a di(lower alkyl)amino-lower alkyl group,
- 25 u) a phenoxy group, unsubstituted or substituted with 1 to

3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group, a lower alkoxy group, a halo-lower alkoxy group and a di(lower alkyl)amino-lower alkyl group,

5 v) a heteroaryl group, unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group, a lower alkoxy group, a halo-lower alkoxy group and a di(lower alkyl)amino-lower alkyl group,

10 w) a heteroaryloxy group, unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group, a lower alkoxy group, a halo-lower alkoxy group and a di(lower alkyl)amino-lower alkyl group, or

15 x) a lower alkoxy group substituted with a group selected from an aryl group or a heteroaryl group, or

when two of X^1 , X^2 , X^3 , X^4 and X^5 are adjacent each other, they are bonded together to form a group represented by $-O(CH_2)_mO-$, $-O(CH_2)_n-$ or $-(CH_2)_p-$;

20 R^{20} and R^{21} are each independently a hydrogen atom, a lower alkyl group, a cycloalkyl group, a heterocycloalkyl group, a bridged cyclichydrocarbon group, a heteroaryl-lower alkyl group, a hydroxy-lower alkyl group, a lower alkoxy-lower alkyl group, a lower acyl group, a lower alkoxycarbonyl group, an
25 aralkyloxycarbonyl group or a di(lower alkyl)amino-lower alkyl group, or

R^{20} and R^{21} , taken together with the nitrogen atom to which

they are bonded, form a cyclic amino group, wherein the cyclic amino group is unsubstituted or substituted with one or two substituents selected independently from the group consisting of:

- 5 a) a lower alkyl group,
- b) a cycloalkyl group,
- c) a phenyl group, unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group,
- 10 a lower alkoxy group and a halo-lower alkoxy group,
- d) an aralkyl group, wherein the ring of the aralkyl group is unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group, a lower alkoxy group
- 15 and a halo-lower alkoxy group, or adjacent ring-carbon atoms of the aralkyl group are substituted with $-O-(CH_2)_m-O-$,
- e) a heteroaryl group,
- f) a heteroaryl-lower alkyl group,
- g) a lower alkyl group substituted with a group selected from
- 20 a hydroxyl group, a lower alkoxy group, a carboxy group, an aralkyloxycarbonyl group, a cyclic aminocarbonyl group or a di(lower alkyl)amino group,
- h) a hydroxyl group,
- i) an oxo group,
- 25 j) a lower alkoxycarbonyl group,
- k) an aralkyloxycarbonyl group,
- l) a carbamoyl group,

- m) a lower acyl group,
- n) a benzoyl group,
- o) a di(lower alkyl)amino group, and
- p) a diphenylmethylene group;

5 A^1 is a bond, a C_{1-3} -alkylene group or a carbonyl group;

A^2 is a bond or a C_{1-3} -alkylene group;

R^{22} is:

- a) a lower alkyl group,
- b) a phenyl group, unsubstituted or substituted with 1 to 3
- 10 substituents selected independently from the group consisting
- of a halogen atom, a lower alkyl group, a halo-lower alkyl group,
- a lower alkoxy group and a halo-lower alkoxy group,
- c) a heteroaryl group, unsubstituted or substituted with 1
- to 3 substituents selected independently from the group
- 15 consisting of a halogen atom, a lower alkyl group, a halo-lower
- alkyl group, a lower alkoxy group and a halo-lower alkoxy group,
- or
- d) a di(lower alkyl)amino-lower alkyl group;

20 R^{23} and R^{24} are each independently a hydrogen atom or a

lower alkyl group, or

R^{23} and R^{24} , taken together with the nitrogen atom to which

they are bonded, form a cyclic amino group, wherein the cyclic

amino group is unsubstituted or substituted with a group selected

from a lower alkyl group or an aralkyl group;

25 m is 1 or 2;

n is 2 or 3;

p is 3 or 4;

X^6 , X^7 and X^8 are each independently:

- a) a halogen atom,
- b) a lower alkyl group,
- c) a halo-lower alkyl group,
- 5 d) a hydroxy-lower alkyl group,
- e) a cycloalkyl group,
- f) a heterocycloalkyl-lower alkyl group,
- g) a lower alkoxy group,
- h) a halo-lower alkoxy group,
- 10 i) a lower acyl group,
- j) a carboxy group,
- k) $-A^1-NR^{20}R^{21}$,
- l) $-A^2-SR^{22}$,
- m) $-SO_2NR^{23}R^{24}$,
- 15 n) a phenyl group, unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group, a lower alkoxy group, a halo-lower alkoxy group and a di(lower alkyl)amino-lower alkyl group,
- 20 o) a phenoxy group, unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group, a lower alkoxy group, a halo-lower alkoxy group and a di(lower alkyl)amino-lower alkyl group,
- 25 p) an aralkyl group, wherein the ring of the aralkyl group is unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a

lower alkyl group, a halo-lower alkyl group, a lower alkoxy group, a halo-lower alkoxy group and a di(lower alkyl)amino-lower alkyl group,

q) a heteroaryl group, unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group, a lower alkoxy group, a halo-lower alkoxy group and a di(lower alkyl)amino-lower alkyl group,

r) a heteroaryloxy group, unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group, a lower alkoxy group, a halo-lower alkoxy group and a di(lower alkyl)amino-lower alkyl group, or

s) an aralkyloxy group;

15 R^3 , R^4 , R^5 and R^6 are each independently:

- a) a hydrogen atom,
- b) a halogen atom,
- c) a lower alkyl group,
- d) a halo-lower alkyl group,
- 20 e) a lower alkoxy group,
- f) a halo-lower alkoxy group,
- g) a hydroxyl group,
- h) a cyano group,
- i) an aryl group, unsubstituted or substituted with 1 to 3
- 25 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group, a lower alkoxy group and a halo-lower alkoxy group,

j) an aralkyloxy group, wherein the ring of the aralkyloxy group is unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group, a lower alkoxy group and a halo-lower alkoxy group,

k) a di(lower alkyl)amino group,

l) a lower alkylsulfanyl group, or

m) a nitro group, or

when two of R^3 , R^4 , R^5 and R^6 are adjacent each other, they are bonded together to form a group represented by $-\text{CH}=\text{CH}-\text{CH}=\text{CH}-$, provided that at least one of R^3 , R^4 , R^5 and R^6 is other than a hydrogen atom;

with the proviso that the following compounds are excluded:

- (1) 1-acetylaminonaphtho[2,1-b]furan-2-carboxamide,
- (2) 1-benzoylaminonaphtho[2,1-b]furan-2-carboxamide,
- (3) 3-benzoylamino-5-chlorobenzofuran-2-carboxamide,
- (4) 5-chloro-3-[2-(3,4-diethoxyphenyl)acetylamino]-benzofuran-2-carboxamide,
- (5) 5-bromo-3-[2-(3,4-diethoxyphenyl)acetylamino]-benzofuran-2-carboxamide,
- (6) 5-chloro-3-(2-chloroacetylamino)benzofuran-2-carboxamide, and
- (7) 3-acetylamino-5-chlorobenzofuran-2-carboxamide.

In another aspect, the present invention provides a pharmaceutical composition which comprises, as an active ingredient, a compound represented by general formula (I) or a pharmaceutically acceptable salt thereof.

In still another aspect, the present invention provides a therapeutic or prophylactic agent for a disease mediated by adenosine A_{2A} receptors, which comprises a compound represented by general formula (I) or a pharmaceutically acceptable salt thereof.

In still another aspect, the present invention provides a pharmaceutical combination comprising a compound represented by general formula (I) or a pharmaceutically acceptable salt thereof and at least one selected from anti-Parkinson drugs, antidepressants, drugs for cognitive function disorders and drugs for cerebral ischemia disorders other than adenosine A_{2A} receptor antagonists.

In still another aspect, the present invention provides a use of a compound represented by general formula (I) or a pharmaceutically acceptable salt thereof for the manufacture of a medicament for treating or preventing a disease mediated by adenosine A_{2A} receptors.

In still another aspect, the present invention provides a method for treating or preventing a disease mediated by adenosine A_{2A} receptors, which comprises administering an effective amount of a compound represented by general formula (I) or a pharmaceutically acceptable salt thereof.

The invention is described using the terms defined below unless otherwise specified.

The term "halogen atom" refers to a fluorine, chlorine, bromine or iodine atom. Preferred halogen atoms for X¹, X², X³,

X^4 and X^5 are a fluorine, chlorine or bromine atom, more preferably a fluorine or chlorine atom and most preferably a fluorine atom. Preferred halogen atoms for X^6 , X^7 and X^8 are a fluorine, chlorine or bromine atom and most preferably a chlorine atom. Preferred
 5 halogen atoms for R^3 are a fluorine atom. Preferred halogen atoms for R^4 are a chlorine or fluorine atom, and more preferably a fluorine atom. Preferred halogen atoms for R^5 and R^6 are a chlorine or fluorine atom.

The term "lower alkyl group" refers to a straight chained
 10 or branched alkyl group having 1 to 6 carbon atoms such as a methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, isopentyl, neopentyl, tert-pentyl, 1-methylbutyl, 2-methylbutyl, 1,2-dimethylpropyl, hexyl, isohexyl group and the like. Preferred lower alkyl groups for
 15 R^1 , R^3 , R^4 , R^5 and R^6 are a C_{1-4} alkyl group, more preferably a methyl, ethyl or isopropyl group, and most preferably a methyl group. Preferred lower alkyl groups for R^2 are a C_{2-4} alkyl group, and more preferably an ethyl, isopropyl or tert-butyl group.

The term "halo-lower alkyl group" refers to a lower alkyl
 20 group substituted with the same or different 1 to 3 halogen atoms such as a fluoromethyl, chloromethyl, bromomethyl, 3-bromopropyl, 4-bromobutyl, 5-bromopentyl, difluoromethyl, trifluoromethyl, 2,2,2-trifluoroethyl group and the like. Preferred halo-lower alkyl groups for R^2 are a 4-bromobutyl or
 25 5-bromopentyl group. Preferred halo-lower alkyl groups for R^4 , R^5 and R^6 are a trifluoromethyl group. Preferred halo-lower alkyl groups for X^1 , X^2 , X^3 , X^4 , X^5 , X^6 , X^7 and X^8 are a chloromethyl

or trifluoromethyl group.

The term "hydroxy-lower alkyl group" refers to a lower alkyl group substituted with a hydroxyl group such as a hydroxymethyl, 1-hydroxyethyl, 2-hydroxyethyl, 2-hydroxy-
 5 2-methylpropyl, 3-hydroxypropyl group and the like.

The term "cycloalkyl group" refers to a 3- to 7-membered saturated cyclic hydrocarbon such as a cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl and cycloheptyl group. Preferred cycloalkyl groups for R² are a cyclopropyl or cyclobutyl group,
 10 and more preferably a cyclopropyl group.

The term "bridged cyclic hydrocarbon group" refers to a 5- to 7-membered bridged saturated cyclic hydrocarbon having 7 to 10 carbon atoms such as a bicyclo[2.2.1]heptane-2-yl, adamantan-1-yl group and the like.

15 The term "cycloalkyloxy group" refers to a group represented by (cycloalkyl)-O- such as a cyclopropyloxy, cyclobutyloxy, cyclopentyloxy, cyclohexyloxy group and the like.

The term "aryl-cycloalkyl group" refers to a cycloalkyl
 20 group substituted with an aryl group such as a 2-phenylcyclopropyl, 2-phenylcyclopentyl, 3-phenylcyclopentyl group and the like.

The term "heterocycloalkyl group" refers to a 4- to 7-membered saturated heterocyclic group having -NH-, -O- or -S-
 25 as a member of the ring such as a tetrahydrofuryl, tetrahydrothienyl, tetrahydropyranyl, pyrrolidin-2-yl, pyrrolidin-3-yl, piperidin-2-yl, piperidin-3-yl,

piperidin-4-yl group and the like. The heterocycloalkyl group may be optionally substituted with one or two lower alkyl or aralkyl groups such as a N-methylpiperidin-4-yl, N-benzylpiperidin-4-yl, N-phenethylpiperidin-4-yl group and
 5 the like.

The term "heterocycloalkyloxy group" refers to a group represented by (heterocycloalkyl)-O- such as a tetrahydropyran-4-yloxy, piperidin-4-yloxy, N-methyl-
 piperidin-4-yloxy, N-benzylpiperidin-4-yloxy, N-phenethyl-
 10 piperidin-4-yloxy group and the like.

The term "heterocycloalkyl-lower alkyl group" refers to a lower alkyl group substituted with a heterocycloalkyl group such as a N-methylpiperidin-4-ylmethyl, N-isopropylpiperidin-4-ylmethyl, N-benzylpiperidin-4-ylmethyl, N-phenethyl-
 15 piperidin-4-ylmethyl group and the like.

The term "alkenyl group" refers to an unsaturated hydrocarbon having 2 to 6 carbon atoms, which may be straight chained or branched and contains at least one double bond such as a vinyl, allyl group and the like.

20 The term "aryl group" refers to an aromatic hydrocarbon group having 6 to 10 carbon atoms such as a phenyl, 1-naphthyl and 2-naphthyl group, preferably a phenyl group.

The term "aralkyl group" refers to a lower alkyl group substituted with an aryl group such as a benzyl, phenethyl,
 25 1-phenylethyl, 3-phenylpropyl, 4-phenylbutyl, naphthylmethyl group and the like, preferably a benzyl, phenethyl, 1-phenylethyl, 3-phenylpropyl or 4-phenylbutyl group.

The term "aryl-alkenyl group" refers to an alkenyl group substituted with an aryl group such as a stylyl, cinnamyl group and the like.

The term "aryloxy group" refers to a group represented
5 by (aryl)-O- such as a phenoxy, 1-naphthyloxy, 2-naphthyloxy group and the like, preferably a phenoxy group.

The term "aralkyloxy group" refers to a group represented
by (aralkyl)-O- such as a benzyloxy, phenethyloxy,
1-phenylethoxy, 3-phenylpropoxy group and the like, preferably
10 a benzyloxy group.

The term "aryloxy-lower alkyl group" refers to a lower
alkyl group substituted with an aryloxy group such as a
phenoxymethyl, 1-phenoxyethyl, 2-phenoxyethyl,
1-methyl-1-phenoxyethyl, 3-phenoxypropyl, naphthyloxymethyl
15 group and the like, preferably a phenoxymethyl, 1-phenoxy-
ethyl, 2-phenoxyethyl or 3-phenoxypropyl group.

The term "aralkyloxy-lower alkyl group" refers to a lower
alkyl group substituted with an aralkyloxy group such as a
benzyloxymethyl, 2-benzyloxyethyl, phenethyloxymethyl,
20 naphthyloxymethyl group and the like, preferably a benzyoxy-
methyl group.

The term "lower alkylsulfanyl group" refers to a group
represented by (lower alkyl)-S- such as a methylsulfanyl,
ethylsulfanyl, propylsulfanyl, isopropylsulfanyl,
25 butylsulfanyl group and the like.

The term "arylsulfanyl group" refers to a group represented
by (aryl)-S- such as a phenylsulfanyl, 1-naphthylsulfanyl,

2-naphthylsulfanyl group and the like, preferably a phenyl-sulfanyl group.

The term "arylsulfanyl-lower alkyl group" refers to a lower alkyl group substituted with an arylsulfanyl group such as a phenylsulfanylmethyl, 1-phenylsulfanylethyl, 2-phenyl-sulfanylethyl, 1-methyl-1-phenylsulfanylethyl, 3-phenyl-sulfanylpropyl, naphthylsulfanylmethyl group and the like, preferably a phenylsulfanylmethyl group.

The term "lower alkoxy group" refers to a straight chained or branched alkoxy group having 1 to 6 carbon atoms such as a methoxy, ethoxy, propoxy, isopropoxy, butoxy isobutoxy, sec-butoxy, tert-butoxy, pentyloxy, hexyloxy group and the like. Preferred lower alkoxy groups for R^3 , R^4 , R^5 , R^6 , X^6 , X^7 and X^8 are a C_{1-3} alkoxy group, more preferably a methoxy or ethoxy group, and most preferably a methoxy group. Preferred alkoxy groups for X^1 , X^2 , X^3 , X^4 and X^5 are a C_{1-4} alkoxy group, and more preferably a methoxy, ethoxy, propoxy or isopropoxy group.

The term "halo-lower alkoxy group" refers to a lower alkoxy group substituted with the same or different 1 to 3 halogen atoms such as a difluoromethoxy, trifluoromethoxy, 2,2,2-trifluoroethoxy group and the like.

The term "lower alkoxy-lower alkyl group" refers to a lower alkyl group substituted with a lower alkoxy group such as a methoxymethyl, 2-methoxyethyl, ethoxymethyl group and the like, preferably a methoxymethyl group.

The term "lower alkoxy-lower alkoxy group" refers to a lower alkoxy group substituted with a lower alkoxy group such

as a 2-methoxyethoxy, 2-ethoxyethoxy, 3-methoxypropoxy, 4-methoxybutoxy group and the like.

The term "lower acyl group" refers to a group represented by H-CO- or (lower alkyl)-CO- such as a formyl, acetyl, propionyl, butyryl, isobutyryl, valeryl, isovaleryl, pivaloyl group and the like, preferably an acetyl group.

The term "lower acyloxy group" refers to a group represented by (lower acyl)-O- such as an acetyloxy, propionyloxy, isobutyryloxy, pivaloyloxy group and the like, preferably an acetyloxy group.

The term "lower alkoxy carbonyl group" refers to a group represented by (lower alkoxy)-CO- such as a methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, isopropoxycarbonyl, butoxycarbonyl, isobutoxycarbonyl, sec-butoxycarbonyl, tert-butoxycarbonyl, pentyloxycarbonyl, hexyloxycarbonyl group and the like.

The term "aralkyloxycarbonyl group" refers to a group represented by (aralkyloxy)-CO- such as a benzyloxycarbonyl group and the like.

The term "di(lower alkyl) amino group" refers to an amino group substituted with two lower alkyl groups such as a dimethylamino, diethylamino, dipropylamino, diisopropylamino group and the like, preferably a dimethylamino or diethylamino group.

The term "di(lower alkyl) amino-lower alkyl group" refers to a lower alkyl group substituted with a di(lower alkyl) amino group such as a dimethylaminomethyl, 2-dimethylaminoethyl,

3-dimethylaminopropyl, diethylaminomethyl, 2-diethylaminoethyl group, preferably a 2-dimethylaminoethyl group.

The term "cyclic amino group" refers to a 5- to 8-membered cyclic amine which may contain -NH-, -O- or -S- as a member of the ring. Examples of cyclic amino group include a 1-pyrrolidyl, piperidino, piperazino, morpholino, thiomorpholino, azepan-1-yl group and the like, preferably a 1-pyrrolidyl, piperidino, morpholino, piperazino or azepan-1-yl group. Adjacent ring carbon atoms of the cyclic amino group may be optionally condensed with a benzene or cycloalkyl ring. Such condensed cyclic amino groups include a indolin-1-yl, 1,2,3,4-tetrahydroquinolin-1-yl, octahydroisoindol-2-yl group and the like. A ring carbon of the cyclic amino group may form a spiro ring together with a 1,2-ethylenedioxy group such as 1,4-dioxa-8-azaspiro[4,5]dec-8-yl group and the like.

The term "cyclic aminocarbonyl group" refers to a group represented by (cyclic amino)-CO- such as pyrrolidinocarbonyl, piperidinocarbonyl, morpholinocarbonyl, piperazinocarbonyl group and the like.

The term "heteroaryl group" refers to a 5- or 6-membered monocyclic aromatic heterocycle having 1 to 5 carbon atoms and 1 to 4 heteroatoms selected independently from the group consisting of an oxygen, nitrogen and sulfur atom, or a 8- to 10-membered bicyclic aromatic heterocycle having 1 to 9 carbon atoms and 1 to 4 heteroatoms selected independently from the group consisting of an oxygen, nitrogen and sulfur atom, provided that said heterocycles do not include adjacent oxygen and/or

sulfur atoms. Examples of monocyclic heteroaryl groups include pyrrolyl, furyl, thienyl, imidazolyl, pyrazolyl, oxazolyl, isoxazolyl, 1,2,4-oxadiazolyl, tetrazolyl, thiazolyl, isothiazolyl, 1,2,3-thiadiazolyl, triazolyl, pyridyl, 5 pyrazinyl, pyrimidyl and pyridazinyl, preferably furyl, thienyl, isoxazolyl or pyridyl, and more preferably furyl. Examples of bicyclic heteroaryl groups include indolyl, indazolyl, benzofuranyl, benzothienyl, benzothiazolyl, quinolyl, isoquinolyl, phthalazinyl, benzimidazolyl, benzoxazolyl and 10 the like, and preferably benzofuranyl. The heterocycles include all position isomers such as 2-pyridyl, 3-pyridyl or 4-pyridyl.

The term "heteroaryl-lower alkyl group" refers to a lower alkyl group substituted with a heteroaryl group such as a 2-furylmethyl, 3-furylmethyl, 2-thienylmethyl, 3-thienyl- 15 methyl, 3-pyridylmethyl, 3-pyridylethyl, 3-benzofurylmethyl, 3-benzothienylmethyl group and the like.

The term " C_{1-3} alkylene group" refers to a bivalent and straight chained saturated hydrocarbon chain having 1 to 3 carbon atoms, which may be optionally substituted with 1 to 3 methyl 20 groups. Examples of C_{1-3} alkylene groups include $-CH_2-$, $-CH_2CH_2-$, $-CH(CH_3)-$, $-CH(CH_3)CH_2-$, $-CH_2CH(CH_3)-$, $-C(CH_3)_2-$, $-C(CH_3)_2CH_2-$, $-CH_2C(CH_3)_2-$, $-CH_2CH_2CH_2-$, $-C(CH_3)_2CH_2CH_2-$, $-C(CH_3)_2CH_2CH(CH_3)-$ and the like, preferably $-CH_2-$, $-CH_2CH_2-$ or $-CH_2CH_2CH_2-$, and more preferably $-CH_2-$.

25 In the case where a compound represented by general formula (I) contains one or more asymmetric carbons, then all stereoisomers in the R- or S-configuration at each of asymmetric

carbons and their mixture are contemplated within the scope of the present invention. In such cases, racemic compounds, racemic mixtures, individual enantiomers and mixtures of diastereomers are also contemplated within the scope of the present invention.

5 In the case where a compound represented by general formula (I) exists in one or more geometrical isomers, then all geometrical isomers such as cis isomer, trans isomer and the mixture thereof are also contemplated within the scope of the present invention. A compound represented by general formula (I) may form a solvate
10 with a pharmaceutically acceptable solvent such as water, ethanol and the like.

Compounds represented by general formula (I) may exist in the form of salts. Examples of such salts include acid addition salts formed with mineral acids such as hydrochloric acid, hydrobromic acid, hydroiodic acid, sulfuric acid, nitric acid,
15 phosphoric acid and the like; acid addition salts formed with organic acids such as formic acid, acetic acid, trifluoroacetic acid, methanesulfonic acid, benzenesulfonic acid, p-toluenesulfonic acid, propionic acid, citric acid, succinic
20 acid, tartaric acid, fumaric acid, butyric acid, oxalic acid, malonic acid, maleic acid, lactic acid, malic acid, carbonic acid, glutamic acid, aspartic acid and the like; basic salts formed with inorganic bases such as sodium, potassium, calcium and the like; basic salts formed with organic bases such as
25 triethylamine, piperidine, morpholine, lysine and the like.

The term "prodrug" as used herein refers to a compound which can be converted into a compound represented by general

formula (I) in vivo. Such prodrugs are also contemplated within the scope of the present invention. Various forms of prodrugs are well known in the art.

In the case where a compound represented by formula (I)
 5 contains a carboxylic acid as a functional group, then a prodrug may include an ester formed by the replacement of the hydrogen atom of the carboxylic acid group with the following groups:
 lower alkyl, lower alkanoyloxymethyl, 1-(lower alkanoyloxy)ethyl, 1-methyl-1-(lower alkanoyloxy)ethyl, lower
 10 alkoxycarbonyloxymethyl, 1-(lower alkoxycarbonyloxy)ethyl, 1-methyl-1-(lower alkoxycarbonyloxy)ethyl, N-(alkoxy-carbonyl)aminomethyl, 1-(N-(lower alkoxycarbonyl)amino)ethyl, 3-phthalidyl, 4-crotonolactonyl, gamma-butyrolacton-4-yl, N,N-di(lower alkyl)amino-lower alkyl such as β -dimethylamino-ethyl, carbamoyl-lower alkyl, N,N-di(lower alkyl)carbamoyl-
 15 lower alkyl, piperidino-lower alkyl, pyrrolidino-lower alkyl or morpholino-lower alkyl.

In the case where a compound represented by formula (I) contains a hydroxyl group, then a prodrug may include a compound
 20 formed by the replacement of the hydrogen atom of the hydroxyl group with the following groups: a lower acyl group such as an acetyl, propionyl, butyryl, isobutyryl, pivaloyl group and the like; a lower alkoxycarbonyl group such as a methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, isopropoxycarbonyl, tert-butoxycarbonyl group and the like; or a succinoyl group.
 25

In the case where a compound represented by formula (I) contains an amino group such as -NH or -NH₂, then a prodrug may

include a compound formed by the replacement of the hydrogen atom of the amino group with the following groups: a lower acyl group such as an acetyl, propionyl, butyryl, isobutyryl, pivaloyl group and the like; or a lower alkoxy-carbonyl group such as a
 5 methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, isopropoxycarbonyl, tert-butoxycarbonyl group and the like.

The prodrug compounds described above may be prepared from compounds represented by general formula (I) according to known methods as described in T.W. Green and P.G.H. Wuts, "Protective
 10 Groups in Organic Synthesis" the third edition and references described therein.

In an embodiment of a compound represented by general formula (I) of the present invention,

15 R^1 is preferably a hydrogen atom;

R^2 is preferably:

a) a lower alkyl group,
 b) a cycloalkyl group,
 c) an aryl group, unsubstituted or substituted with 1 to 5
 20 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 ,

d) an aralkyl group, wherein the ring of the aralkyl group is unsubstituted or substituted with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 ,

25 e) a lower alkoxy-lower alkyl group,

f) an aryloxy-lower alkyl group, wherein the ring of the aryloxy-lower alkyl group is unsubstituted or substituted with

1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 ,

g) an aralkyloxy-lower alkyl group, wherein the ring of the aralkyloxy-lower alkyl group is unsubstituted or substituted
5 with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 , or

h) a heteroaryl group, unsubstituted or substituted with 1 to 3 substituents selected from the group consisting of X^6 , X^7 and X^8 ; and

10 R^2 is more preferably:

a) a cycloalkyl group,

b) an aryl group, unsubstituted or substituted with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 , or

15 c) a heteroaryl group, unsubstituted or substituted with 1 to 3 substituents selected from the group consisting of X^6 , X^7 and X^8 ;

preferred X^1 , X^2 , X^3 , X^4 and X^5 are each independently:

a) a halogen atom,

20 b) a lower alkyl group,

c) a lower alkoxy group,

d) a halo-lower alkoxy group,

e) a heterocycloalkyloxy group,

f) a hydroxyl group,

25 g) $-A^1-NR^{20}R^{21}$,

h) $-A^2-SR^{22}$,

i) $-SO_2NR^{23}R^{24}$,

j) a heteroaryl group, unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group, a lower alkoxy group, a halo-lower alkoxy group and di(lower alkyl)amino-lower alkyl group, or

when two of X^1 , X^2 , X^3 , X^4 and X^5 are adjacent each other, they are bonded together to form $-OCH_2O-$;

preferred X^6 , X^7 and X^8 are each independently:

- a) a halogen atom,
- 10 b) a lower alkyl group,
- c) a hydroxy-lower alkyl group,
- d) a cycloalkyl group,
- e) a heterocycloalkyl-lower alkyl group,
- f) $-A^1-NR^{20}R^{21}$,
- 15 g) $-SO_2NR^{23}R^{24}$,
- h) a phenyl group, unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group, a lower alkoxy group, a halo-lower alkoxy group and a di(lower
- 20 alkyl)amino-lower alkyl group, or

i) a phenoxy group, unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group, a lower alkoxy group, a halo-lower alkoxy group and a di(lower

25 alkyl)amino-lower alkyl group;

R^3 is preferably a hydrogen atom; and

preferred R^4 , R^5 and R^6 are each independently:

- a) a hydrogen atom,
- b) a halogen atom,
- c) a lower alkyl group,
- d) a halo-lower alkyl group, or
- 5 e) a lower alkoxy group, provided that at least one of R^4 , R^5 and R^6 is other than a hydrogen atom.

In a preferable embodiment of the present invention,
 R^1 is a hydrogen atom.

- 10 In a more preferable embodiment of the present invention,
 R^1 is a hydrogen atom, and
 R^3 is a hydrogen atom.

In an even more preferable embodiment of the present invention,

- 15 R^1 is a hydrogen atom,
 R^2 is:

- a) a lower alkyl group,
- b) a cycloalkyl group,
- c) an aryl group, unsubstituted or substituted with 1 to 5
- 20 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 ,

d) an aralkyl group, wherein the ring of the aralkyl group is unsubstituted or substituted with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 ,

- 25 e) a lower alkoxy-lower alkyl group,
- f) an aryloxy-lower alkyl group, wherein the ring of the aryloxy-lower alkyl group is unsubstituted or substituted with

1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 ,

g) an aralkyloxy-lower alkyl group, wherein the ring of the aralkyloxy-lower alkyl group is unsubstituted or substituted
5 with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 , or

h) a heteroaryl group, unsubstituted or substituted with 1 to 3 substituents selected from the group consisting of X^6 , X^7 and X^8 , and

10 R^3 is a hydrogen atom.

In an even more preferable embodiment of the present invention,

R^1 is a hydrogen atom,

R^2 is:

15 a) a lower alkyl group,

b) a cycloalkyl group,

c) an aryl group, unsubstituted or substituted with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 ,

20 d) an aralkyl group, wherein the ring of the aralkyl group is unsubstituted or substituted with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 ,

e) a lower alkoxy-lower alkyl group,

f) an aryloxy-lower alkyl group, wherein the ring of the
25 aryloxy-lower alkyl group is unsubstituted or substituted with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 ,

g) an aralkyloxy-lower alkyl group, wherein the ring of the aralkyloxy-lower alkyl group is unsubstituted or substituted with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 , or

- 5 h) a heteroaryl group, unsubstituted or substituted with 1 to 3 substituents selected from the group consisting of X^6 , X^7 and X^8 ,

R^3 is a hydrogen atom, and

R^4 , R^5 and R^6 are each independently:

- 10 a) a hydrogen atom,
 b) a halogen atom,
 c) a lower alkyl group,
 d) a halo-lower alkyl group, or
 e) a lower alkoxy group, provided that at least one of R^4 ,
 15 R^5 and R^6 is other than a hydrogen atom.

In a still more preferable embodiment of the present invention,

R^1 is a hydrogen atom,

R^2 is:

- 20 a) a cycloalkyl group,
 b) an aryl group, unsubstituted or substituted with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 , or
 c) a heteroaryl group, unsubstituted or substituted with 1
 25 to 3 substituents selected from the group consisting of X^6 , X^7 and X^8 ,

R^3 is a hydrogen atom, and

R^4 , R^5 and R^6 are each independently:

- a) a hydrogen atom,
- b) a halogen atom,
- c) a lower alkyl group,
- 5 d) a halo-lower alkyl group, or
- e) a lower alkoxy group, provided that at least one of R^4 , R^5 and R^6 is other than a hydrogen atom.

In a still more preferable embodiment of the present invention,

10 R^1 is a hydrogen atom,

R^2 is:

- a) a cycloalkyl group,
- b) an aryl group, unsubstituted or substituted with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 ,
- 15 X^4 and X^5 , or

- c) a heteroaryl group, unsubstituted or substituted with 1 to 3 substituents selected from the group consisting of X^6 , X^7 and X^8 ,

R^3 is a hydrogen atom,

20 R^4 , R^5 and R^6 are each independently:

- a) a hydrogen atom,
- b) a halogen atom,
- c) a lower alkyl group,
- d) a halo-lower alkyl group, or
- 25 e) a lower alkoxy group, provided that at least one of R^4 , R^5 and R^6 is other than a hydrogen atom,

X^1 , X^2 , X^3 , X^4 and X^5 are each independently:

- a) a halogen atom,
- b) a lower alkyl group,
- c) a lower alkoxy group,
- d) a halo-lower alkoxy group,
- 5 e) a heterocycloalkyloxy group,
- f) a hydroxyl group,
- g) $-A^1-NR^{20}R^{21}$,
- h) $-A^2-SR^{22}$,
- i) $-SO_2NR^{23}R^{24}$,
- 10 j) a heteroaryl group, unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group, a lower alkoxy group, a halo-lower alkoxy group and di(lower alkyl)amino-lower alkyl group, or
- 15 when two of X^1 , X^2 , X^3 , X^4 and X^5 are adjacent each other, they are bonded together to form $-OCH_2O-$; and
- X^6 , X^7 and X^8 are each independently:
 - a) a halogen atom,
 - b) a lower alkyl group,
 - 20 c) a hydroxy-lower alkyl group,
 - d) a cycloalkyl group,
 - e) a heterocycloalkyl-lower alkyl group,
 - f) $-A^1-NR^{20}R^{21}$,
 - g) $-SO_2NR^{23}R^{24}$,
 - 25 h) a phenyl group, unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group,

a lower alkoxy group, a halo-lower alkoxy group and a di(lower alkyl)amino-lower alkyl group, or

i) a phenoxy group, unsubstituted or substituted with 1 to 3 substituents selected independently from the group consisting of a halogen atom, a lower alkyl group, a halo-lower alkyl group, a lower alkoxy group, a halo-lower alkoxy group and a di(lower alkyl)amino-lower alkyl group.

Specific examples of preferred embodiments of the present invention are compounds selected from the group consisting of:

- 10 (1) 3-cyclopropanecarbonylamino-5-fluorobenzofuran-2-carboxamide;
- (2) 5-chloro-3-cyclopropanecarbonylamino-2-carboxamide;
- (3) 3-(3-fluorobenzoylamino)-6-methoxybenzofuran-2-
- 15 carboxamide;
- (4) 3-(4-fluorobenzoylamino)-6-methoxybenzofuran-2-carboxamide;
- (5) 5-fluoro-3-(3-methylbenzoylamino)benzofuran-2-carboxamide;
- 20 (6) 3-(benzo[1,3]dioxole-5-carbonyl)amino-6-fluorobenzofuran-2-carboxamide;
- (7) 5-chloro-3-(furan-2-carbonyl)aminobenzofuran-2-carboxamide;
- (8) 5,7-difluoro-3-(furan-2-carbonyl)aminobenzofuran-2-
- 25 carboxamide;
- (9) 5,7-difluoro-3-(5-methylfuran-2-carbonyl)aminobenzofuran-2-carboxamide;

- (10) 3-(5-ethylfuran-2-carbonyl)amino-5-fluorobenzofuran-2-carboxamide;
- (11) 3-(5-ethylfuran-2-carbonyl)amino-5,7-difluorobenzofuran-2-carboxamide;
- 5 (12) 6-methoxy-3-(5-phenylfuran-2-carbonyl)aminobenzofuran-2-carboxamide;
- (13) 6-fluoro-3-(6-phenoxy pyridine-3-carbonyl)aminobenzofuran-2-carboxamide;
- (14) 6-methoxy-3-(2-methoxyacetyl amino)benzofuran-2-
- 10 carboxamide;
- (15) 3-[2-(4-chlorophenoxy)acetyl amino]-5-fluorobenzofuran-2-carboxamide;
- (16) 3-(2-benzyloxyacetyl amino)-5-fluorobenzofuran-2-carboxamide;
- 15 (17) 6-chloro-3-cyclopropanecarbonylaminobenzofuran-2-carboxamide;
- (18) 3-cyclopropanecarbonyl amino-5,7-difluorobenzofuran-2-carboxamide;
- (19) 7-chloro-3-cyclopropanecarbonyl amino-5-fluoro-
- 20 benzofuran-2-carboxamide;
- (20) 3-cyclopropanecarbonyl amino-5-fluoro-7-methoxybenzofuran-2-carboxamide;
- (21) 3-cyclobutanecarbonyl amino-5,7-difluorobenzofuran-2-carboxamide;
- 25 (22) 5-fluoro-7-methoxy-3-(4-methoxybenzoyl amino)-benzofuran-2-carboxamide;
- (23) 5,7-difluoro-3-phenylacetylaminobenzofuran-2-

carboxamide;

(24) 5,7-difluoro-3-[3-(4-methylpiperazine-1-carbonyl)-
benzoylamino]benzofuran-2-carboxamide;

(25) 6-methoxy-3-[3-(4-phenylpiperazin-1-ylmethyl)benzoyl-
5 amino]benzofuran-2-carboxamide;

(26) 6-methoxy-3-[4-(1-methyl-1H-imidazol-2-ylsulfanyl-
methyl)benzoylamino]benzofuran-2-carboxamide;

(27) 3-[5-(4-benzylpiperazin-1-ylmethyl)furan-2-carbonyl]-
amino-5,7-difluorobenzofuran-2-carboxamide;

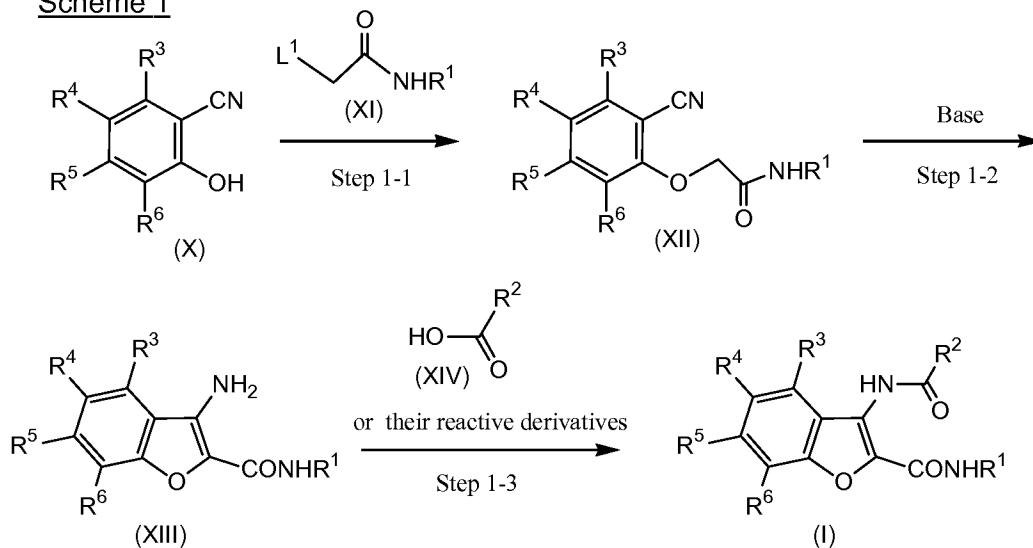
10 (28) 3-[5-(4-benzo[1,3]dioxol-5-ylmethylpiperazin-1-
ylmethyl)furan-2-carbonyl]amino-5,7-difluorobenzofuran-2-
carboxamide;

(29) tert-butyl 4-[5-(2-carbamoyl-5,7-difluorobenzofuran-3-
ylcarbamoyl)furan-2-ylmethyl]piperazine-1-carboxylate, and

15 (30) 5-fluoro-3-[5-(1-hydroxyethyl)furan-2-carbonyl]amino-
benzofuran-2-carboxamide,

or a pharmaceutically acceptable salt thereof.

A compound represented by general formula (I) can be
20 prepared by methods as illustrated in schemes 1 to 4.

Scheme 1

wherein R^1, R^2, R^3, R^4, R^5 and R^6 are as defined above, L^1 is a leaving group such as a chlorine, bromine or iodine atom, a methanesulfonyloxy, p-toluenesulfonyloxy group or the like. (Step 1-1)

Condensation of 2-hydroxybenzonitrile derivative (X) with compound (XI) in the presence of a base in an inert solvent provides compound (XII). The solvents employed in the condensation reaction include ethanol, acetonitrile, N,N-dimethylformamide, dimethylsulfoxide, tetrahydrofuran, 1-methyl-2-pyrrolidone or their mixed solvents. The bases include potassium carbonate, sodium carbonate, cesium carbonate, triethylamine, N,N-diisopropylethylamine or the like. The reaction is carried out ordinarily at 0°C to room temperature. The reaction time varies depending on the starting materials employed, the solvent, the reaction temperature or the like, but is usually 1 to 24 hours.

(Step 1-2)

Thereafter, cyclization of the compound (XII) in the presence of a base in an inert solvent provides compound (XIII). The solvents employed in the reaction include ethanol, acetonitrile, N,N-dimethylformamide, dimethylsulfoxide, 5 tetrahydrofuran, 1-methyl-2-pyrrolidone or their mixed solvents. The bases include potassium carbonate, sodium carbonate, cesium carbonate, triethylamine, N,N-diisopropylethylamine or the like. The reaction is carried out ordinarily at room temperature to reflux temperature. The 10 reaction time varies depending on the starting materials employed, the solvent, the reaction temperature or the like, but is usually 1 to 24 hours.

Alternatively, the compound (XIII) can be prepared from compound (X) by conducting steps 1-1 and 1-2 without isolating 15 compound (XII).
(Step 1-3)

The compound (XIII) is condensed with carboxylic acid (XIV) in the presence of a condensing agent such as dicyclohexylcarbodiimide, 1-(3-dimethylaminopropyl)-3- 20 ethylcarbodiimide hydrochloride, ethyl cyanophosphate, diphenylphosphoryl azide or the like in an inert solvent to afford compound (I).

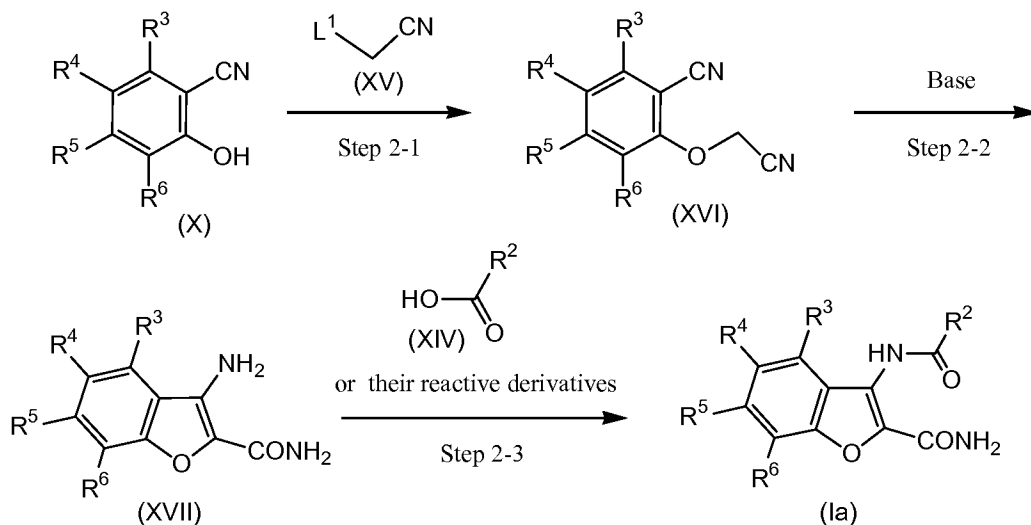
Alternatively, the compound (I) can be prepared by converting carboxylic acid (XIV) into their reactive derivatives 25 such as acid halide, acid anhydride, mixed acid anhydride, 4-nitrophenyl ester, 2,5-dioxapyrrolidine ester or the like, followed by condensing with compound (XIII) in the presence or

absence of a base. Solvents employed in the reaction include acetonitrile, N,N-dimethylformamide, tetrahydrofuran, methylene chloride or their mixed solvents. The bases include potassium carbonate, triethylamine, N,N-diisopropylethylamine, pyridine, N-methylmorpholine, N,N-dimethylaniline or the like. The reaction is carried out ordinarily at -20°C to reflux temperature. The reaction time varies depending on the starting materials employed, the solvent, the reaction temperature or the like, but is usually 15 minutes to 24 hours.

10

Among compounds represented by general formula (I), compound (Ia) wherein R^1 is a hydrogen atom can be also prepared by methods as illustrated in scheme 2.

Scheme 2



wherein R^2 , R^3 , R^4 , R^5 , R^6 and L^1 are as defined above.

15 (Step 2-1)

Condensation of 2-hydroxybenzonitrile derivative (X)

with compound (XV) in the presence of a base in an inert solvent provides compound (XVI). The solvents employed in the condensation reaction include ethanol, acetonitrile, 1-methyl-2-pyrrolidone, N,N-dimethylformamide or the like. The bases include potassium carbonate, sodium carbonate, cesium carbonate, triethylamine, N,N-diisopropylethylamine or the like. The reaction is carried out ordinarily at 0°C to 80°C. The reaction time varies depending on the starting materials employed, the solvent, the reaction temperature or the like, but is usually 1 to 24 hours.

(Step 2-2)

Thereafter, the compound (XVI) is cyclized of in the presence of a base in an inert solvent to provide compound (XVII). The solvents employed in the reaction include alcohols such as methanol, ethanol, isopropanol or the like, and a mixed solvent of the alcohols and water. The bases include potassium hydroxide, sodium hydroxide or the like. The reaction is carried out ordinarily at room temperature to reflux temperature. The reaction time varies depending on the starting materials employed, the solvent, the reaction temperature or the like, but is usually 1 to 24 hours.

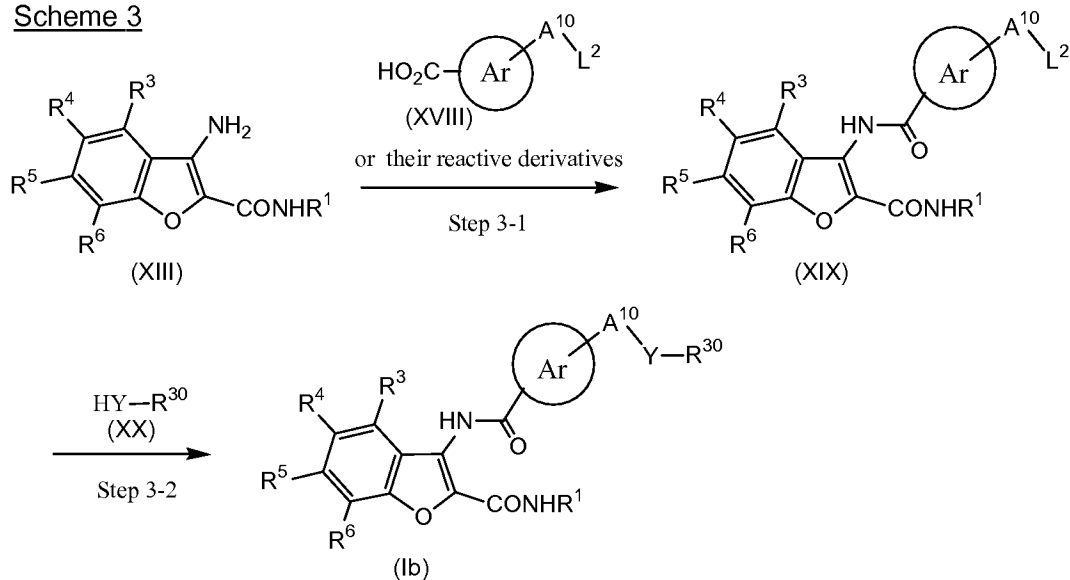
(Step 2-3)

The compound (XVII) is then condensed with carboxylic acid (XIV) according to procedures analogous to those as described in step 1-3 to afford compound (Ia).

Among compounds represented by general formula (I),

compound (Ib) can be prepared by methods as illustrated in scheme 3.

Scheme 3



wherein $\text{R}^1, \text{R}^3, \text{R}^4, \text{R}^5$ and R^6 are as defined above; A^{10} is a C_{1-3} alkylene group; Ar is an aryl or heteroaryl group; L^2 is a leaving group such as a chlorine, bromine or iodine atom, a methanesulfonyloxy, p-toluenesulfonyloxy group or the like; Y is $-\text{N}(\text{R}^{20})-$ or $-\text{S}-$; when Y is $-\text{N}(\text{R}^{20})-$, R^{30} is the same as R^{21} , and when Y is $-\text{S}-$, R^{30} is the same as R^{22} .

(Step 3-1)

Compound (XIII) is condensed with compound (XVIII) according to procedures analogous to those as described in step 1-3 to afford compound (XIX).

(Step 3-2)

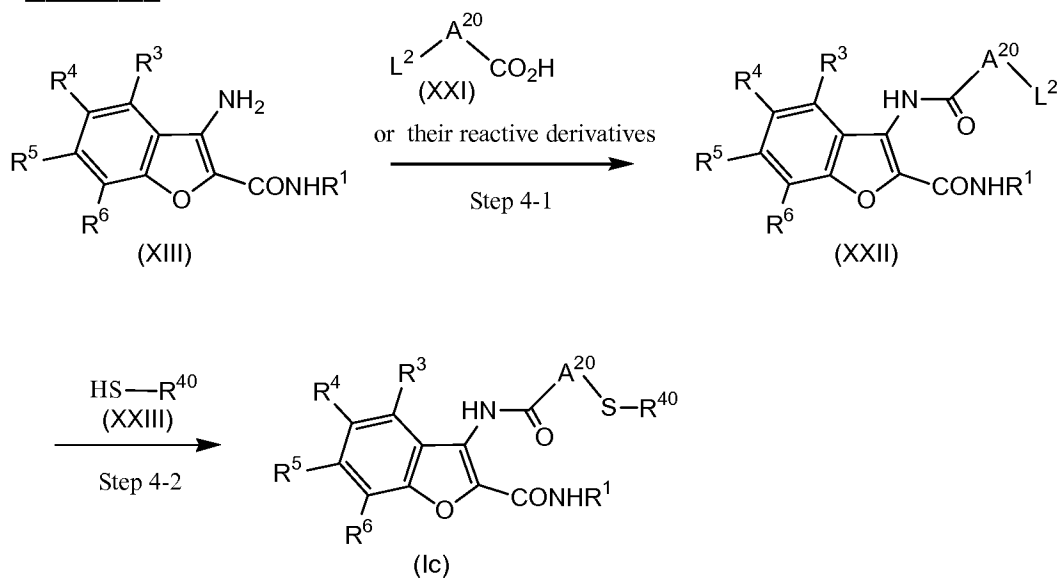
Thereafter, the compound (XIX) is treated with compound (XX) in the presence or absence of a base in an inert solvent to afford compound (Ib). The solvents employed in the reaction include ethanol, isopropanol, acetonitrile,

N,N-dimethylformamide, tetrahydrofuran, methylene chloride, 1-methyl-2-pyrrolidone or their mixed solvents. The bases include potassium carbonate, triethylamine, N,N-diisopropylethylamine, pyridine, N-methylmorpholine, N,N-dimethylaniline or the like. The reaction is carried out ordinarily at -20°C to reflux temperature. The reaction time varies depending on the starting materials employed, the solvent, the reaction temperature or the like, but is usually 15 minutes to 24 hours.

10

Among compounds represented by general formula (I), compound (Ic) can be prepared by methods as illustrated in scheme 4.

Scheme 4



wherein R^1 , R^3 , R^4 , R^5 , R^6 and L^2 are as defined above; A^{20} is a C_{1-6} alkylene group; R^{40} is an aryl group, unsubstituted or substituted with 1 to 5 substituents selected from the group consisting of X^1 , X^2 , X^3 , X^4 and X^5 ; and X^1 , X^2 , X^3 , X^4 and X^5 are

15

as defined above.

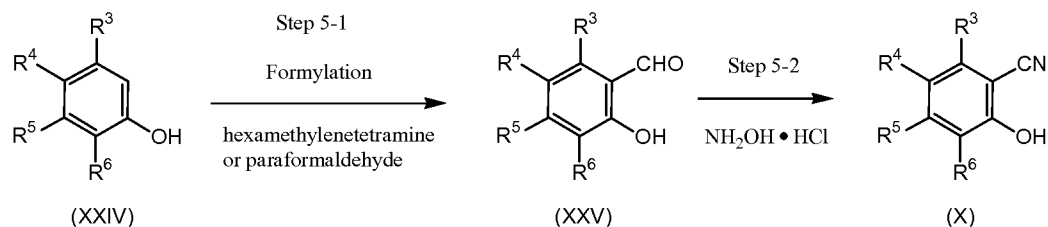
(Step 4-1)

Compound (XIII) is condensed with compound (XXI) according to procedures analogous to those as described in step 1-3 to
5 afford compound (XXII).

(Step 4-2)

Thereafter, the compound (XXII) is treated with compound (XXIII) in the presence of a base in an inert solvent to afford compound (Ic). The solvents employed in the reaction include
10 ethanol, isopropanol, acetonitrile, N,N-dimethylformamide, tetrahydrofuran, methylene chloride, 1-methyl-2-pyrrolidone, or their mixed solvents. The bases include potassium carbonate, triethylamine, N,N-diisopropylethylamine, pyridine, N-methylmorpholine, N,N-dimethylaniline or the like. The
15 reaction is carried out ordinarily at -20°C to reflux temperature. The reaction time varies depending on the starting materials employed, the solvent, the reaction temperature or the like, but is usually 15 minutes to 24 hours.

20 Among the starting materials employed in scheme 1, compound (X) can be prepared by methods as illustrated in scheme 5 or 6.

Scheme 5

wherein R^3 , R^4 , R^5 and R^6 are as defined above.

(Step 5-1)

Phenol derivative (XXIV) is condensed with hexamethylenetetramine in a solvent of trifluoroacetic acid to afford an imine compound. Hydrolysis of the imine compound using an acid provides compound (XXV). The condensation reaction is carried out ordinarily at room temperature to reflux temperature, and the reaction time varies depending on the starting materials employed, the solvent, the reaction temperature or the like, but is usually 1 to 144 hours. The acids employed in the hydrolysis reaction include an aqueous solution of sulfuric acid or the like. The hydrolysis reaction is carried out ordinarily at room temperature to reflux temperature, and the reaction time is usually 15 minutes to 12 hours.

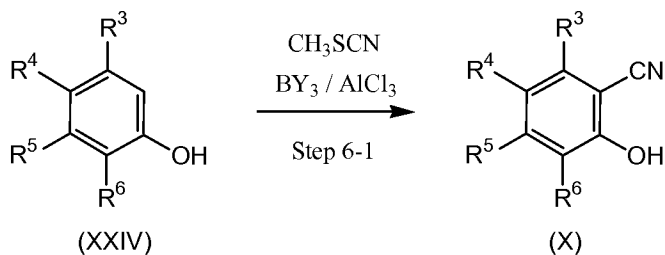
Alternatively, the compound (XXV) can be prepared by condensing phenol derivative (XXIV) with paraformaldehyde in the presence of magnesium chloride and a base in an inert solvent such as acetonitrile or the like. The bases employed in the reaction include triethylamine, *N,N*-diisopropylethylamine or the like. The condensation reaction is carried out ordinarily at room temperature to reflux temperature. The reaction time varies depending on the starting materials employed, the solvent,

the reaction temperature or the like, but is usually 0.5 to 48 hours.

(Step 5-2)

Thereafter, the compound (XXV) is treated with
 5 hydroxylamine hydrochloride in a suitable solvent such as
 1-methyl-2-pyrrolidone, formic acid or the like to afford
 compound (X). The reaction is carried out ordinarily at room
 temperature to reflux temperature. The reaction time varies
 depending on the starting materials employed, the solvent, the
 10 reaction temperature or the like, but is usually 1 to 144 hours.

Scheme 6



wherein R³, R⁴, R⁵ and R⁶ are as defined above; and Y is a chlorine or bromine atom.

(Step 6-1)

15 Compound (X) can be also prepared by treating phenol
 derivative (XXIV) with methylthiocyanic acid in the presence
 of Lewis acid in an inert solvent. The solvents employed in the
 reaction include dichloromethane, 1,2-dichloroethane, or the
 like. Lewis acids include boron trihalides such as boron
 20 trichloride or boron tribromide/aluminum chloride or the like.
 The reaction is carried out ordinarily at 0°C to reflux

temperature. The reaction time varies depending on the starting materials employed, the solvent, the reaction temperature or the like, but is usually 1 to 24 hours.

5 The forementioned schemes are exemplary for preparing compounds represented by general formula (I) of the present invention and synthetic intermediates thereof. Those skilled in the art will appreciate that various changes or modifications of the forementioned schemes may be made without departing from
10 the scope of the invention.

 Compounds represented by general formula (I) of the present invention and intermediates for preparing the compounds of the present invention can be isolated or purified, if required, according to conventional isolation or purification techniques
15 well known to those in the art, such as solvent extraction, crystallization, recrystallization, chromatography, preparative high performance liquid chromatography or the like.

 Compounds of general formula (I) prepared in the
20 above-mentioned schemes exhibit excellent adenosine A_{2A} receptor antagonistic activities, and are accordingly useful as a therapeutic or prophylactic agent for motor function disorders such as Parkinson's disease, Huntington's disease or Wilson's disease and the like; depression; an anxiety disorder; cognitive
25 function disorders such as Alzheimer disease and the like; cerebral ischemia disorders such as cerebral apoplexy, brain lesion after cerebral vasospasm and the like; restless legs

syndrome and the like.

Compounds of general formula (I) can be used, if required, in combination with anti-Parkinson drugs, antidepressants, drugs for cognitive function disorders and drugs for cerebral
 5 ischemia disorders other than adenosine A_{2A} receptor antagonists.

Examples of anti-Parkinson drugs which may be used in combination with compounds of general formula (I) include levodopa, levodopa/carbidopa, levodopa/benserazide, droxidopa,
 10 melevodopa, threodops; dopamine D_2 receptor agonists such as cabergoline, bromocriptine mesylate, terguride, talipexole hydrochloride, ropinirole hydrochloride, pergolide mesylate, pramipexole hydrochloride, rotigotine and the like; anticholinergic agents such as profenamine, trihexyphenidyl
 15 hydrochloride, mazaticol hydrochloride, biperiden, piroheptine hydrochloride, methixene hydrochloride and the like; COMT (catechol O-methyl transferase) inhibitors such as tolcapone, entacapone and the like; NMDA antagonists such as budipine and the like; monoamine oxidase B inhibitors such as selegiline
 20 hydrochloride, rasagiline mesylate and the like; zonisamide; amantadine hydrochloride and the like.

Examples of antidepressants which may be used in combination with compounds of general formula (I) include selective serotonin reuptake inhibitors such as fluoxetine
 25 hydrochloride, sertraline hydrochloride, paroxetine hydrochloride, citalopram hydrobromide, fluvoxamine maleate and the like; selective noradrenalin reuptake inhibitors such

as desipramine hydrochloride, amitriptyline hydrochloride, nortriptyline hydrochloride, reboxetine and the like; serotonin/noradrenalin reuptake inhibitors such as venlafaxine hydrochloride, bupropion hydrochloride, nefazodone
 5 hydrochloride, milnacipran hydrochloride and the like.

Examples of drugs for cognitive function disorders which may be used in combination with compounds of general formula (I) include acetylcholine esterase inhibitors such as tacrine, donepezil hydrochloride, rivastigmine tartrate, metrifonate,
 10 galantamine hydrobromide; memantine hydrochloride; aripiprazole; S-8510; AC-3933 and the like.

Examples of drugs for cerebral ischemia disorders which may be used in combination with compounds of general formula (I) include thrombolytic agents such as t-PA (tissue plasminogen
 15 activator), urokinase and the like; trombin inhibitors such as argatroban and the like; TXA₂ synthase inhibitors such as ozagrel sodium and the like; radical scavengers such as ebselen, edaravone, nicaraven and the like; 5-HT_{1A} agonists such as SUN-N4057, BAYx3702 and the like; NMDA antagonists such as
 20 aptiganel hydrochloride and the like; AMPA antagonists such as S-1746 and the like; Rho kinase inhibitors such as fasudil and the like; src inhibitors and the like.

Various dosage forms of pharmaceutical compositions comprising a compound represented by general formula (I) or a
 25 pharmaceutically acceptable salt thereof, can be administered depending on their usages. Exemplary dosage forms include powders, granules, fine granules, dry syrups, tablets, capsules,

injections, liquids, ointments, suppositories, poultices and the like, which are administered orally or parenterally.

Pharmaceutical compositions can be formulated by admixing, diluting or dissolving with appropriate pharmaceutical additives such as excipients, disintegrators, binders, lubricants, diluents, buffers, isotonic agents, preservatives, wetting agents, emulsifying agents, dispersing agents, stabilizing agents, solubilizing agents and the like, according to a conventional formulation procedure depending upon their dosage forms.

The dosage of a compound represented by general formula (I) or a pharmaceutically acceptable salt thereof is appropriately determined depending on the age, sex or body weight of the individual patient, the severity of the disease, the condition to be treated and the like. A typical dosage for oral administration is in the range of from about 1mg to about 5000 mg per day per adult human. A typical dosage for parenteral administration is in the range of from about 0.1mg to about 500 mg per day per adult human. The dosages may be administered in single or divided doses, for example one to several times daily.

A pharmaceutical combination comprising a compound represented by general formula (I) or a pharmaceutically acceptable salt thereof, and at least one selected from anti-Parkinson drugs, antidepressants, drugs for cognitive function disorders and drugs for cerebral ischemia disorders other than adenosine A_{2A} receptor antagonists, can be administered as a single pharmaceutical composition comprising

all of active ingredients, or as separately formulated pharmaceutical compositions each of which comprises a single active ingredient. Where separately formulated pharmaceutical compositions are used, the compositions may be administered
5 separately, concurrently or at different intervals. Alternatively, where separately formulated pharmaceutical compositions are used, the compositions may be mixed together with an appropriate diluent, and administered simultaneously.

In a pharmaceutical combination comprising a compound
10 represented by general formula (I) or a pharmaceutically acceptable salt thereof, and at least one selected from anti-Parkinson drugs, antidepressants, drugs for cognitive function disorders and drugs for cerebral ischemia disorders other than adenosine A_{2A} receptor antagonists, the dosage of
15 each active ingredient may be appropriately determined depending on the age, sex or body weight of the individual patient, the severity of the disease, administration time, dosage form, administration method, combination of active ingredients and the like.

20 Compounds represented by general formula (I) of the present invention exhibit potent adenosine A_{2A} receptor antagonistic activities. Moreover, preferable compounds of the present invention show selective antagonistic activities on adenosine A_{2A} receptors. Therefore, compounds of the present invention
25 are useful for treating or preventing diseases mediated by adenosine A_{2A} receptors such as motor function disorders, depression, an anxiety disorder, cognitive function disorders,

cerebral ischemia disorder, restless legs syndrome and the like, and are especially useful for treating or preventing Parkinson's disease.

5 **BEST MODE FOR CARRYING OUT THE INVENTION**

The following reference examples, examples and test examples illustrate the invention in further detail. It is to be understood, however, that they are not to be construed as limiting the scope of the invention in any way.

10

Reference Example 1-1

3-Bromo-5-fluoro-2-hydroxybenzaldehyde

To a solution of 2-bromo-4-fluorophenol (2g) in trifluoroacetic acid (10mL) was added hexamethylenetetramine
15 (2.94g) at room temperature, and the resulting mixture was stirred under reflux for 20 hours. A 50% aqueous solution of sulfuric acid was added to the reaction mixture, and the mixture was stirred for further 4 hours at room temperature. The reaction mixture was extracted with ethyl acetate, and washed with water,
20 1mol/L hydrochloric acid and brine. The organic layer was dried over anhydrous magnesium sulfate, and concentrated under reduced pressure. The residue was purified by column chromatography on silica gel to give the title compound (1.93g).

¹H-NMR (CDCl₃) δ ppm: 7.25-7.30 (1H, m), 7.56-7.61 (1H, m), 9.83
25 (1H, s), 11.35 (1H, s)

Reference Examples 1-2 and 1-3 were prepared in a manner

similar to those as described in Reference Example 1-1 using 2,4-difluorophenol and 4-fluoro-2-methoxyphenol instead of 2-bromo-4-fluorophenol.

Reference Example 1-2

5 3,5-Difluoro-2-hydroxybenzaldehyde

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 7.08-7.20 (2H, m), 9.88 (1H, d, $J=1.9\text{Hz}$), 10.72 (1H, s)

Reference Example 1-3

5-Fluoro-2-hydroxy-3-methoxybenzaldehyde

10 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 3.92 (3H, s), 6.85-6.90 (2H, m), 9.87 (1H, s), 10.80 (1H, brs)

Reference Example 2-1

2-Fluoro-6-hydroxybenzonitrile

15 Hydroxylamine hydrochloride (0.60g) was added to a solution of 2-fluoro-6-hydroxybenzaldehyde (1g) in 1-methyl-2-pyrrolidone (10ml), and the resulting mixture was stirred at 120°C for 2days. The reaction mixture was extracted with ethyl acetate, washed with water and brine. The organic layer was dried
20 over anhydrous magnesium sulfate, and concentrated under reduced pressure. The residue was purified by column chromatography on silica gel to give the title compound (0.35g).

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 6.30-6.86 (3H, m), 7.36-7.50 (1H, m)

25 Reference Examples 2-2 to 2-20 were prepared in a manner similar to those as described in Reference Example 2-1 using the corresponding 2-hydroxybenzaldehydes instead of 2-fluoro-

6-hydroxybenzaldehyde.

Reference example 2-2

5-Fluoro-2-hydroxybenzonitrile

¹H-NMR (DMSO-d₆) δ ppm: 6.95-7.05 (1H, m), 7.30-7.45 (1H, m),
5 7.50-7.65 (1H, m), 11.05 (1H, s)

Reference example 2-3

4-Fluoro-2-hydroxybenzonitrile

¹H-NMR (CDCl₃) δ ppm: 6.20-6.35 (1H, m), 6.65-6.80 (2H, m),
7.45-7.60 (1H, m)

10 Reference example 2-4

3,5-Difluoro-2-hydroxybenzonitrile

¹H-NMR (DMSO-d₆) δ ppm: 7.46-7.74 (2H, m), 11.42 (1H, brs)

Reference example 2-5

3-Bromo-5-fluoro-2-hydroxybenzonitrile

15 ¹H-NMR (DMSO-d₆) δ ppm: 7.69-7.76 (1H, m), 7.89-7.96 (1H, m),
11.01 (1H, brs)

Reference example 2-6

2-Hydroxy-5-methylbenzonitrile

¹H-NMR (CDCl₃) δ ppm: 2.29 (3H, s), 6.89 (1H, d, J=8.4Hz),
20 7.24-7.32 (2H, m)

Reference example 2-7

2-Hydroxy-3-methylbenzonitrile

¹H-NMR (CDCl₃) δ ppm: 2.28 (3H, s), 5.87 (1H, brs), 6.86-6.94
(1H, m), 7.30-7.40 (2H, m)

25 Reference example 2-8

1-Hydroxynaphthalene-2-carbonitrile

¹H-NMR (CDCl₃) δ ppm: 7.35-7.85 (5H, m), 8.21-8.38 (1H, m)

Reference example 2-9

2-Hydroxy-5-methoxybenzonitrile

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 3.78 (3H, s), 5.77 (1H, brs), 6.88-6.98 (2H, m), 7.02-7.08 (1H, m)

5 Reference example 2-10

2-Hydroxy-4-methoxybenzonitrile

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 3.94 (3H, s), 6.27 (1H, brs), 6.86-7.16 (3H, m)

Reference example 2-11

10 4-Benzyloxy-2-hydroxybenzonitrile

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 5.08 (2H, s), 6.54-6.64 (2H, m), 7.30-7.48 (6H, m)

Reference example 2-12

2-Hydroxy-3-methoxybenzonitrile

15 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 3.83 (3H, s), 6.46-6.58 (2H, m), 7.40 (1H, d, $J=8.8\text{Hz}$)

Reference example 2-13

3-Ethoxy-2-hydroxybenzonitrile

20 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.48 (3H, t, $J=6.9\text{Hz}$), 4.15 (2H, q, $J=6.9\text{Hz}$), 6.27 (1H, s), 6.80-7.15 (3H, m)

Reference example 2-14

4-Diethylamino-2-hydroxybenzonitrile

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.06-1.13 (6H, m), 3.28-3.37 (4H, m), 6.13-6.26 (2H, m), 7.25 (1H, d, $J=8.8\text{Hz}$), 10.35 (1H, s)

25 Reference example 2-15

4,5-Difluoro-2-hydroxybenzonitrile

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 6.90-7.00 (1H, m), 7.85-7.95 (1H, m),

11.57 (1H, brs)

Reference example 2-16

3-Chloro-5-fluoro-2-hydroxybenzonitrile

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 7.18-7.25 (1H, m), 7.34-7.39 (1H, m)

5 Reference example 2-17

5-Fluoro-2-hydroxy-3-methoxybenzonitrile

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 3.94 (3H, s), 6.04 (1H, s), 6.75-6.85 (2H, m)

Reference example 2-18

10 4-Ethyl-2-hydroxybenzonitrile

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.25 (3H, t, $J=7.6\text{Hz}$), 2.67 (2H, q, $J=7.6\text{Hz}$), 6.78-6.92 (2H, m), 7.40-7.50 (1H, m)

Reference example 2-19

2-Hydroxy-4-isopropylbenzonitrile

15 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.24 (6H, d, $J=6.9\text{Hz}$), 2.75-3.00 (1H, m), 6.35 (1H, brs), 6.80-6.95 (2H, m), 7.30-7.50 (1H, m)

Reference example 2-20

4-Ethoxy-2-hydroxybenzonitrile

20 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.42 (3H, t, $J=6.9\text{Hz}$), 4.04 (2H, q, $J=6.9\text{Hz}$), 6.40-6.60 (2H, m), 7.30-7.45 (1H, m)

Reference example 3-1

4-Chloro-2-hydroxybenzonitrile

To a solution of 3-chlorophenol (2.86mL) in dichloroethane
 25 (48mL) were added aluminum chloride (4.00g) and boron tribromide (3.39mL) under ice-cooling, and methyl thiocyanate (2.46mL) was added to the reaction mixture, and stirred at room temperature

until aluminum chloride was dissolved. Then, the reaction mixture was stirred and heated at 120°C for 20 hours. After cooling, a 4mol/L aqueous solution of sodium hydroxide (99mL) was added to the reaction mixture, and the mixture was stirred at
 5 approximately 80°C for 30 minutes. The solution was washed with dichloromethane. The aqueous layer was acidified with 6mol/L hydrochloric acid, and extracted with diethyl ether. The organic layer was washed with brine and dried over anhydrous magnesium sulfate. The organic solvent was concentrated under reduced
 10 pressure. The residue was treated with hexane and collected by filtration to give the title compound (2.5g).

¹H-NMR (CDCl₃) δ ppm: 6.99 (1H, dd, J=1.8, 8.5Hz), 7.03 (1H, d, J=1.8Hz), 7.44 (1H, d, J=8.5Hz)

15 Reference Example 3-2

3-Hydroxybiphenyl-4-carbonitrile

Reference Example 3-2 was prepared in a manner similar to those as described in Reference Example 3-1 using 3-hydroxybiphenyl instead of 3-chlorophenol.

20 ¹H-NMR (CDCl₃) δ ppm: 6.10 (1H, brs), 7.18-7.28 (2H, m), 7.39-7.50 (3H, m), 7.54-7.60 (3H, m)

Reference example 4-1

3-Amino-4-fluorobenzofuran-2-carboxamide

25 To a solution of 2-fluoro-6-hydroxybenzonitrile (0.35g) in ethanol (20mL) were added potassium carbonate (0.54g) and bromoacetoamide (0.43g) successively at room temperature, and

the resulting mixture was heated under reflux for 2 hours. Then, potassium hydroxide (0.29g) was added to the reaction mixture and heated under reflux for 12 hours. The reaction mixture was diluted with water, and the organic solvent was concentrated under reduced pressure. The resulting precipitate was collected by filtration to give the title compound (0.25g).

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 5.24 (2H, brs), 5.81 (2H, brs), 6.80-6.96 (1H, m), 7.10-7.42 (2H, m)

Reference Examples 4-2 to 4-27 were prepared in a manner similar to those as described in Reference Example 4-1 using the corresponding 2-hydroxybenzonitriles instead of 2-fluoro-6-hydroxybenzonitrile. These were illustrated in table 1.

Table 1

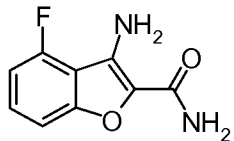
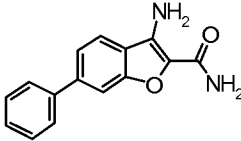
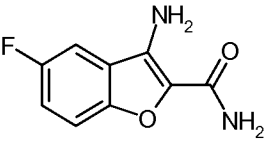
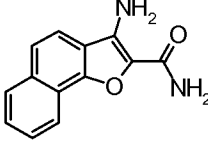
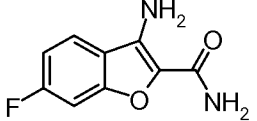
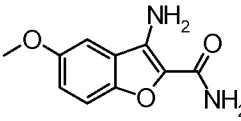
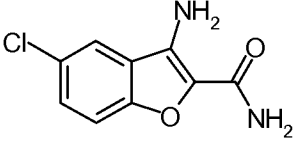
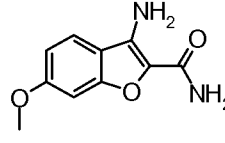
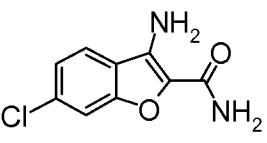
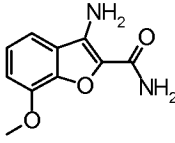
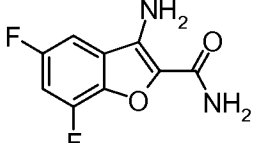
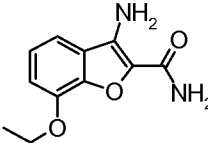
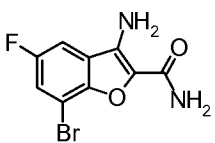
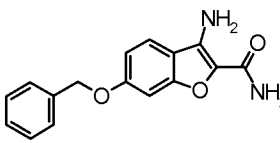
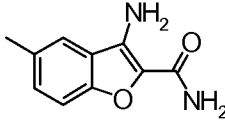
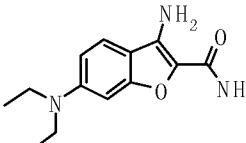
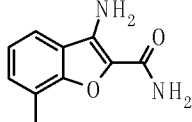
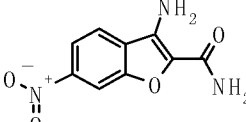
Reference example	Structure	Reference example	Structure
4-1		4-10	
4-2		4-11	
4-3		4-12	
4-4		4-13	
4-5		4-14	
4-6		4-15	
4-7		4-16	
4-8		4-17	
4-9		4-18	

Table 1 (continued)

Reference example	Structure	Reference example	Structure
4-19		4-24	
4-20		4-25	
4-21		4-26	
4-22		4-27	
4-23			

The physical data of reference examples 4-2 to 4-27 were shown as below.

5 Reference example 4-2

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 5.97 (2H, s), 7.08-7.72 (5H, m)

Reference example 4-3

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 5.03 (2H, brs), 5.71 (2H, brs), 6.96-7.14 (2H, m), 7.44-7.54 (1H, m)

10 Reference example 4-4

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 6.02 (2H, s), 7.12-7.50 (4H, m),

7.94-8.00 (1H, m)

Reference example 4-5

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 6.07 (2H, brs), 7.10-7.36 (3H, m),
7.48-7.58 (1H, m), 7.80-7.92 (1H, m)

5 Reference example 4-6

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 6.07 (2H, brs), 7.15-7.65 (4H, m)

Reference example 4-7

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 6.06 (2H, brs), 7.29 (2H, s), 7.62-7.78
(2H, m)

10 Reference example 4-8

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 2.39 (3H, s), 5.92 (2H, s), 7.04-7.32
(4H, m), 7.57-7.65 (1H, m)

Reference example 4-9

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 2.44 (3H, s), 5.93 (2H, brs), 7.06-7.28
15 (4H, m), 7.63 (1H, d, $J=7.6\text{Hz}$)

Reference example 4-10

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 6.05 (2H, brs), 7.22 (2H, brs), 7.35-7.80
(7H, m), 7.92 (1H, d, $J=8.2\text{Hz}$)

Reference example 4-11

20 $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 6.05 (2H, brs), 7.30 (2H, brs), 7.52-8.40
(6H, m)

Reference example 4-12

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 3.79 (3H, s), 5.90 (2H, s), 7.02 (1H,
dd, $J=2.7, 9.0\text{Hz}$), 7.17 (2H, brs), 7.31 (1H, d, $J=9.0\text{Hz}$), 7.40
25 (1H, d, $J=2.7\text{Hz}$)

Reference example 4-13

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 3.93 (3H, s), 5.98 (2H, s), 6.97-7.44

(5H, m)

Reference example 4-14

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 3.81 (3H, s), 5.97 (2H, s), 6.83-6.95 (2H, m), 7.04 (2H, brs), 7.71 (1H, d, $J=8.7\text{Hz}$)

5 Reference example 4-15

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.52 (3H, t, $J=6.9\text{Hz}$), 4.24 (2H, q, $J=6.9\text{Hz}$), 4.99 (2H, brs), 6.85-7.00 (1H, m), 7.05-7.20 (2H, m)

Reference example 4-16

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 5.18 (2H, s), 5.99 (2H, s), 6.91-7.02 (2H, m), 7.08 (2H, brs), 7.30-7.52 (5H, m), 7.73 (1H, d, $J=8.7\text{Hz}$)

10

Reference example 4-17

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 1.05-1.18 (6H, m), 3.28-3.52 (4H, m), 5.87 (2H, brs), 6.47 (1H, s), 6.60-6.72 (1H, m), 6.84 (2H, brs), 7.55 (1H, d, $J=8.9\text{Hz}$)

15 Reference example 4-18

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 6.21 (2H, s), 7.51 (2H, brs), 8.08-8.26 (3H, m)

Reference example 4-19

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 6.04 (2H, brs), 7.29 (2H, brs), 7.55-7.70 (1H, m), 7.85-7.95 (1H, m)

20

Reference example 4-20

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 3.94 (3H, s), 5.93 (2H, brs), 6.93-7.04 (1H, m), 7.13-7.45 (3H, m)

Reference example 4-21

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 2.31-2.36 (3H, m), 5.96 (2H, s), 7.21 (2H, brs), 7.30-7.37 (1H, m), 7.55-7.65 (1H, m)

25

Reference example 4-22

¹H-NMR (DMSO-d₆) δ ppm: 2.42 (3H, s), 5.98 (2H, s), 6.95-7.35 (4H, m), 7.68-7.74 (1H, m)

Reference example 4-23

¹H-NMR (CDCl₃) δ ppm: 1.28 (3H, t, J=7.6Hz), 2.77 (2H, q, J=7.6Hz),
5 4.99 (2H, brs), 7.06-7.22 (2H, m), 7.40-7.48 (1H, m)

Reference example 4-24

¹H-NMR (CDCl₃) δ ppm: 1.30 (6H, d, J=6.9Hz), 2.90-3.15 (1H, m),
4.99 (2H, brs), 5.71 (2H, brs), 7.05-7.30 (2H, m), 7.40-7.50 (1H, m)

10 Reference example 4-25

¹H-NMR (CDCl₃) δ ppm: 1.37 (9H, s), 4.99 (2H, brs), 7.28-7.50 (3H, m)

Reference example 4-26

¹H-NMR (DMSO-d₆) δ ppm: 6.16 (2H, s), 7.41 (2H, brs), 7.55-7.65
15 (1H, m), 7.73-7.80 (1H, m), 8.05-8.15 (1H, m)

Reference example 4-27

¹H-NMR (CDCl₃) δ ppm: 1.46 (3H, t, J=6.9Hz), 4.07 (2H, q, J=6.9Hz),
4.99 (2H, brs), 6.80-6.90 (2H, m), 7.35-7.45 (1H, m)

20 Reference example 5-1

4-Phenoxybenzoyl chloride

To a suspension of 4-phenoxybenzoic acid (1.4g) in dichloromethane (15mL) was added N,N-dimethylformamide (0.05mL), and oxalyl chloride (1.2mL) was added dropwise to the
25 mixture under ice-cooling. The resulting mixture was stirred at room temperature for 1 hour, and the reaction mixture was concentrated under reduced pressure to give the title compound

(1.54g).

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 6.95-7.15 (4H, m), 7.20-7.30 (1H, m),
7.40-7.50 (2H, m), 8.05-8.15 (2H, m)

5 Reference examples 5-2 and 5-3 were prepared in a manner
similar to those as described in reference example 5-1 using
the corresponding carboxylic acids instead of 4-phenoxybenzoic
acid.

Reference example 5-2

10 5-Ethylfuran-2-carbonyl chloride

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.30 (3H, t, $J=7.5\text{Hz}$), 2.76 (2H, q,
 $J=7.5\text{Hz}$), 6.20-6.30 (1H, m), 7.40-7.45 (1H, m)

Reference example 5-3

3-Bromo-4-fluorobenzoyl chloride

15 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 7.20-7.30 (1H, m), 8.05-8.15 (1H, m),
8.30-8.40 (1H, m)

Reference example 6-1

2-Carbamoyl-5-fluoro-7-hydroxybenzofuran-3-yl ammonium

20 bromide

To a mixture of 3-amino-5-fluoro-7-methoxybenzofuran-
2-carboxamide (4.44g) and dichloromethane (100mL) was added a
1 mol/L solution of boron tribromide in dichloromethane (100mL)
under ice-cooling, and the resulting mixture was stirred at room
25 temperature overnight. The reaction mixture was poured into
methanol (200mL) under ice-cooling, and the mixture was stirred
for another 1 hour. The solvent of the mixture was removed under

reduced pressure. To the residue was added methanol, and the resulting precipitate was collected by filtration to give the title compound (4.0g).

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 6.62-6.70 (1H, m), 7.00-7.30 (3H, m)

5

Reference example 6-2

2-Carbamoyl-7-hydroxybenzofuran-3-yl ammonium bromide

Reference Example 6-2 was prepared in a manner similar to those as described in Reference Example 6-1 using 3-amino-7-methoxybenzofuran-2-carboxamide instead of 3-amino-5-fluoro-7-methoxybenzofuran-2-carboxamide.

10

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 6.60-7.50 (3H, m), 7.61 (1H, d, $J=8.5\text{Hz}$)

Reference example 7-1

15 3-Amino-7-chloro-5-fluorobenzofuran-2-carboxamide

To a mixture of 3-chloro-5-fluoro-2-hydroxybenzonitrile (24.8g), potassium carbonate (60.1g) and 1-methyl-2-pyrrolidone (150mL) was added chloroacetonitrile (11.0mL) at room temperature, and the resulting mixture was stirred at 60°C for 5 hours. The reaction mixture was poured into ice/water, and the precipitate was collected by filtration. The solid was dissolved with ethyl acetate (250mL) and purified by column chromatography on aminopropylated silica gel (eluent: ethyl acetate) to give 3-chloro-2-cyanomethoxy-5-fluorobenzonitrile (23.88g).

20

25

To a solution of 3-chloro-2-cyanomethoxy-5-fluorobenzonitrile in ethanol (200mL) was added potassium hydroxide

(9.5g), and the mixture was stirred under reflux overnight. Water (750mL) was added to the reaction mixture, and the resulting precipitate was collected by filtration. The solid was suspended in ethyl acetate (500mL) and purified by column chromatography on aminopropylated silica gel (eluent: ethyl acetate) to give the title compound (13.09g).

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 6.09 (2H, s), 7.34 (2H, brs), 7.55-7.61 (1H, m), 7.67-7.74 (1H, m)

10 Reference example 8-1

Methyl 5-[2-(4-methoxyphenyl)ethyl]furan-2-carboxylate

Sodium hydride (60%, 0.09g) was added to a mixture of (5-methoxycarbonylfuran-2-ylmethyl) triphenylphosphonium chloride (0.823g), 4-methoxybenzaldehyde (0.257g) and

15 N,N-dimethylformamide (10mL), and the resulting mixture was stirred at room temperature for 3 hours. Water and ethyl acetate were added to the reaction mixture, and the organic layer was separated. The organic layer was washed with 1mol/L hydrochloric acid, a 1mol/L aqueous solution of sodium hydroxide, 1mol/L hydrochloric acid, water and brine successively, and dried over
20 anhydrous magnesium sulfate. The solvent was removed under reduced pressure, and the residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate = 8/1) to give methyl 5-[2-(4-methoxyphenyl)vinyl]furan-2-
25 carboxylate.

A mixture of methyl 5-[2-(4-methoxyphenyl)vinyl]furan-2-carboxylate, 10% palladium on carbon (50% wet, 0.01g) and ethyl

acetate (30mL) was stirred at room temperature under hydrogen atmosphere for 1.5 hours. After the catalyst was removed by filtration, the solvent was removed under reduced pressure to give the title compound (0.344g).

5 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 2.90-3.02 (4H, m), 3.79 (3H, s), 3.89 (3H, s), 6.07 (1H, d, $J=3.5\text{Hz}$), 6.78-6.88 (2H, m), 7.05-7.15 (3H, m)

Reference examples 8-2 to 8-6 were prepared in a manner similar to those as described in Reference Example 8-1 using the corresponding aldehydes or ketones instead of 4-methoxybenzaldehyde.

Reference example 8-2

Methyl 5-[2-(3-methoxyphenyl)ethyl]furan-2-carboxylate

15 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 2.92-3.05 (4H, m), 3.79 (3H, s), 3.89 (3H, s), 6.10 (1H, d, $J=3.5\text{Hz}$), 6.70-6.82 (3H, m), 7.06-7.24 (2H, m)

Reference example 8-3

Methyl 5-[2-(2-methoxyphenyl)ethyl]furan-2-carboxylate

20 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 2.99 (4H, s), 3.82 (3H, s), 3.89 (3H, s), 6.09 (1H, d, $J=3.5\text{Hz}$), 6.81-6.91 (2H, m), 7.05-7.24 (3H, m)

Reference example 8-4

Methyl 5-(1-isopropylpiperidin-4-ylmethyl)furan-2-carboxylate

25 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.02 (6H, d, $J=6.6\text{Hz}$), 1.20-1.38 (2H, m), 1.62-1.77 (3H, m), 2.00-2.15 (2H, m), 2.56-2.90 (5H, m), 3.87 (3H, s), 6.12 (1H, d, $J=3.5\text{Hz}$), 7.09 (1H, d, $J=3.5\text{Hz}$)

Reference example 8-5

Methyl 5-(1-benzylpiperidin-4-ylmethyl)furan-2-carboxylate

¹H-NMR (CDCl₃) δ ppm: 1.22-1.38 (2H, m), 1.58-1.80 (3H, m),
 1.87-2.00 (2H, m), 2.61 (2H, d, J=7.3Hz), 2.80-2.90 (2H, m),
 5 3.48 (2H, s), 3.87 (3H, s), 6.11 (1H, d, J=3.5Hz), 7.09 (1H,
 d, J=3.5Hz), 7.20-7.40 (5H, m)

Reference example 8-6

Methyl

5-(1-phenethylpiperidin-4-ylmethyl)furan-2-carboxylate

10 MS (ESI, m/z) : 328 (M+H)+

Reference example 9-1

5-[2-(4-Methoxyphenyl)ethyl]furan-2-carboxylic acid

To a solution of methyl 5-[2-(4-methoxyphenyl)ethyl]-
 15 furan-2-carboxylate (0.344g) in ethanol (5mL) was added a 2mol/L
 aqueous solution of sodium hydroxide (1.35mL), and the resulting
 mixture was heated under reflux for 2 hours. To the reaction
 mixture were added water and diethyl ether, and the aqueous layer
 was separated. Then, the aqueous layer was acidified with 1mol/L
 20 hydrochloric acid (6mL). The acidified aqueous layer was
 extracted with ethyl acetate. The organic layer was washed with
 brine, and then dried over anhydrous magnesium sulfate. The
 solvent was removed under reduced pressure to give the title
 compound (0.296g).

25 ¹H-NMR (CDCl₃) δ ppm: 2.90-3.07 (4H, m), 3.79 (3H, s), 6.12 (1H,
 d, J=3.5Hz), 6.80-6.86 (2H, m), 7.06-7.14 (2H, m), 7.23 (1H,
 d, J=3.5Hz)

Reference Examples 9-2 to 9-26 were prepared in a manner similar to those as described in Reference Example 9-1 using the corresponding carboxylate ester instead of methyl

5 5-[2-(4-methoxyphenyl)ethyl]furan-2-carboxylate.

Reference example 9-2

5-[2-(3-Methoxyphenyl)ethyl]furan-2-carboxylic acid

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 2.95-3.10 (4H, m), 3.79 (3H, s), 6.15 (1H, d, $J=3.6\text{Hz}$), 6.70-6.80 (3H, m), 7.18-7.25 (2H, m)

10 Reference example 9-3

5-[2-(2-Methoxyphenyl)ethyl]furan-2-carboxylic acid

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 3.01 (4H, s), 3.82 (3H, s), 6.14 (1H, d, $J=3.4\text{Hz}$), 6.80-6.93 (2H, m), 7.05-7.25 (3H, m)

Reference example 9-4

15 5-(1-Isopropylpiperidin-4-ylmethyl)furan-2-carboxylic acid

MS (ESI, m/z): 252 (M+H)+

Reference example 9-5

5-(1-Benzylpiperidin-4-ylmethyl)furan-2-carboxylic acid

MS (ESI, m/z): 300 (M+H)+

20 Reference example 9-6

5-(1-Phenethylpiperidin-4-ylmethyl)furan-2-carboxylic acid

MS (ESI, m/z): 314 (M+H)+

Reference example 9-7

5-(Azepane-1-sulfonyl)furan-2-carboxylic acid

25 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.55-1.85 (8H, m), 3.40-3.50 (4H, m), 7.00-7.10 (1H, m), 7.30-7.35 (1H, m)

Reference example 9-8

5-(4-Methylpiperazine-1-sulfonyl)furan-2-carboxylic acid

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.26 (3H, s), 2.45-2.60 (4H, m), 3.05-3.20 (4H, m), 7.05-7.10 (1H, m), 7.20-7.25 (1H, m)

Reference example 9-9

5 3-Cyclohexyloxybenzoic acid

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.25-1.65 (6H, m), 1.75-2.10 (4H, m), 4.25-4.40 (1H, m), 7.10-7.20 (1H, m), 7.30-7.45 (1H, m), 7.55-7.75 (2H, m)

Reference example 9-10

10 3-(Tetrahydropyran-4-yloxy)benzoic acid

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.75-1.90 (2H, m), 1.95-2.10 (2H, m), 3.50-3.70 (2H, m), 3.95-4.05 (2H, m), 4.50-4.65 (1H, m), 7.10-7.25 (1H, m), 7.30-7.45 (1H, m), 7.60-7.75 (2H, m)

Reference example 9-11

15 4-Benzyloxy-3-morpholin-4-ylmethylbenzoic acid

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.25-2.50 (4H, m), 3.40-3.70 (4H, m), 3.59 (2H, brs), 5.22 (2H, s), 7.10-7.55 (6H, m), 7.80-8.00 (2H, m), 12.66 (1H, brs)

Reference example 9-12

20 4-Ethoxy-3-morpholin-4-ylmethylbenzoic acid hydrochloride

$^1\text{H-NMR}$ ($\text{CDCl}_3 + \text{MeOD-d}_4$) δ ppm: 1.50 (3H, t, $J=7.0\text{Hz}$), 2.85-3.05 (2H, m), 3.35-3.45 (2H, m), 3.90-4.00 (1H, m), 4.15-4.25 (4H, m), 4.34 (2H, s), 6.95-7.05 (1H, m), 8.10-8.25 (2H, m)

Reference example 9-13

25 6-(2-Fluorophenoxy)nicotinic acid

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 7.00-7.15 (1H, m), 7.15-7.30 (4H, m), 8.30-8.40 (1H, m), 8.86 (1H, d, $J=2.1\text{Hz}$)

Reference example 9-14

6-(3-Fluorophenoxy)nicotinic acid

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 6.90-7.05 (4H, m), 7.35-7.45 (1H, m),
8.30-8.40 (1H, m), 8.90 (1H, d, $J=2.2\text{Hz}$)

5 Reference example 9-15

6-(4-Fluorophenoxy)nicotinic acid

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 6.99 (1H, d, $J=8.4\text{Hz}$), 7.05-7.20 (4H, m),
8.30-8.40 (1H, m), 8.88 (1H, d, $J=2.5\text{Hz}$)

Reference example 9-16

10 6-(2,4-Difluorophenoxy)nicotinic acid

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 6.85-7.05 (2H, m), 7.08 (1H, d, $J=9.0\text{Hz}$),
7.15-7.30 (1H, m), 8.30-8.45 (1H, m), 8.83 (1H, d, $J=2.2\text{Hz}$)

Reference example 9-17

6-(2,4,6-Trifluorophenoxy)nicotinic acid

15 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 6.75-6.90 (2H, m), 7.10-7.20 (1H, m),
8.35-8.45 (1H, m), 8.80-8.85 (1H, m)

Reference example 9-18

6-(2-Isopropylphenoxy)nicotinic acid

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.19 (6H, d, $J=7.0\text{Hz}$), 3.00-3.15 (1H, m),
20 6.94 (1H, d, $J=8.9\text{Hz}$), 7.00-7.10 (1H, m), 7.20-7.30 (2H, m),
7.35-7.45 (1H, m), 8.25-8.35 (1H, m), 8.91 (1H, d, $J=2.0\text{Hz}$)

Reference example 9-19

6-(4-Isopropylphenoxy)nicotinic acid

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.27 (6H, d, $J=6.9\text{Hz}$), 2.85-3.00 (1H, m),
25 6.90-7.00 (1H, m), 7.05-7.15 (2H, m), 7.20-7.35 (2H, m),
8.25-8.35 (1H, m), 8.90-8.95 (1H, m)

Reference example 9-20

6-(4-Fluoro-2-methylphenoxy)nicotinic acid

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 2.15 (3H, s), 6.90-7.10 (4H, m), 8.30-8.40 (1H, m), 8.87 (1H, d, $J=2.0\text{Hz}$)

Reference example 9-21

5 6-(2-Methoxyphenoxy)nicotinic acid

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 3.76 (3H, s), 6.95-7.10 (3H, m), 7.10-7.20 (1H, m), 7.20-7.30 (1H, m), 8.25-8.35 (1H, m), 8.87 (1H, d, $J=1.9\text{Hz}$)

Reference example 9-22

10 6-(3-Methoxyphenoxy)nicotinic acid

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 3.81 (3H, s), 6.65-6.85 (3H, m), 6.90-7.00 (1H, m), 7.34 (1H, t, $J=8.2\text{Hz}$), 8.25-8.40 (1H, m), 8.92 (1H, d, $J=2.3\text{Hz}$)

Reference example 9-23

15 6-(4-Fluoro-2-methoxyphenoxy)nicotinic acid

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 3.74 (3H, s), 6.65-6.80 (2H, m), 7.00 (1H, d, $J=9.0\text{Hz}$), 7.05-7.15 (1H, m), 8.25-8.35 (1H, m), 8.85 (1H, d, $J=2.1\text{Hz}$)

Reference example 9-24

20 6-(Pyridin-3-yloxy)nicotinic acid

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 7.33 (1H, d, $J=8.4\text{Hz}$), 7.85-7.95 (1H, m), 8.15-8.30 (1H, m), 8.30-8.45 (1H, m), 8.65-8.75 (2H, m), 8.87 (1H, d, $J=2.1\text{Hz}$)

Reference example 9-25

25 6-(3-Hydroxymethylphenoxy)nicotinic acid

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 4.52 (2H, brs), 5.25 (1H, brs), 7.00-7.25 (4H, m), 7.35-7.45 (1H, m), 8.25-8.35 (1H, m), 8.60-8.70 (1H,

m), 13.17 (1H, brs)

Reference example 9-26

6-Phenethyloxynicotinic acid

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 3.11 (2H, t, $J=7.1\text{Hz}$), 4.62 (2H, t, $J=7.1\text{Hz}$),

5 6.70-6.85 (1H, m), 7.15-7.40 (5H, m), 8.15-8.25 (1H, m),

8.85-8.95 (1H, m)

Reference example 10-1

Benzyl 3-(4-methylpiperazine-1-carbonyl)benzoate

10 To a solution of 1-methylpiperazine (0.656mL) in
dichloromethane (5mL) was added a solution of benzyl
3-chlorocarbonylbenzoate (2.14g) in dichloromethane (5mL) and
the resulting mixture was stirred at room temperature overnight.
To the reaction mixture were added water and ethyl acetate, and
15 the organic layer was separated. The organic layer was washed
with a 1mol/L aqueous solution of sodium hydroxide, water and
brine successively, and dried over anhydrous sodium sulfate.
The solvent was removed under reduced pressure, and the residue
was purified by column chromatography on aminopropylated silica
20 gel (eluent: ethyl acetate) to give the title compound (1.7g).

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 2.24-2.60 (7H, m), 3.41 (2H, brs), 3.81
(2H, brs), 5.37 (2H, s), 7.30-7.64 (7H, m), 8.08-8.16 (2H, m)

Reference Example 10-2

25 Benzyl 3-(morpholin-4-carbonyl)benzoate

Reference example 10-2 was prepared in a manner similar
to those as described in reference example 10-1 using morpholine

instead of 1-methylpiperazine.

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 3.30-3.95 (8H, m), 5.38 (2H, s), 7.31-7.65 (7H, m), 8.07-8.18 (2H, m)

5 Reference example 11-1

3-(4-Methylpiperazine-1-carbonyl)benzoic acid

A mixture of benzyl 3-(4-methylpiperazine-1-carbonyl)-benzoate (1.7g) and 10% palladium on carbon (50%wet, 0.128g) in ethanol (50mL) was stirred at room temperature under hydrogen
10 atmosphere for 3 days. To the reaction mixture was added a mixed solvent of trifluoroethanol (50mL) and dichloromethane (50ml), and the mixture was stirred at room temperature for 10 minutes. The catalyst was removed by filtration, and the filtrate was concentrated under reduced pressure to give the title compound
15 (1.15g).

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.16-2.46 (7H, m), 3.00-3.80 (4H, m), 7.50-7.66 (2H, m), 7.83-8.06 (2H, m)

Reference example 11-2

20 3-(Morpholine-4-carbonyl)benzoic acid

Reference example 11-2 was prepared in a manner similar to those as described in reference example 11-1 using benzyl 3-(morpholine-4-carbonyl)benzoate instead of benzyl 3-(4-methylpiperazine-1-carbonyl)benzoate.

25 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.20-3.80 (8H, m), 7.55-7.70 (2H, m), 7.90-8.04 (2H, m), 13.13 (1H, brs)

Reference example 12-1

Methyl 5-(4-methylpiperazine-1-sulfonyl)furan-2-carboxylate

To a solution of methyl 5-chlorosulfonyl-2-furan-carboxylate (0.414g) in tetrahydrofuran (5mL) was added
 5 1-methylpiperazine (0.41mL) under ice-cooling, and the resulting mixture was stirred at room temperature for 1 day. To the reaction mixture were added a saturated aqueous solution of sodium carbonate and ethyl acetate. The organic layer was separated, washed with water and brine, and dried over anhydrous
 10 magnesium sulfate. The solvent was removed under reduced pressure to give the title compound (0.265g).

¹H-NMR (CDCl₃) δ ppm: 2.30 (3H, s), 2.40-2.55 (4H, m), 3.20-3.40 (4H, m), 3.92 (3H, s), 7.04 (1H, d, J=3.5Hz), 7.20 (1H, d, J=3.5Hz)

15 Reference Examples 12-2 to 12-4 were prepared in a manner similar to those as described in Reference Example 12-1 using the corresponding sulfonyl chlorides and amines instead of methyl 5-chlorosulfonyl-2-furan carboxylate and 1-methylpiperazine.

Reference example 12-2

20 Methyl 5-(azepane-1-sulfonyl)furan-2-carboxylate

¹H-NMR (CDCl₃) δ ppm: 1.55-1.80 (8H, m), 3.35-3.45 (4H, m), 3.92 (3H, s), 6.95-7.05 (1H, m), 7.15-7.20 (1H, m)

Reference example 12-3

3-(4-Methylpiperazine-1-sulfonyl)benzoic acid hydrochloride

25 ¹H-NMR (DMSO-d₆) δ ppm: 2.73 (3H, s), 2.95-4.10 (8H, m), 7.75-7.95 (1H, m), 8.00-8.10 (1H, m), 8.15-8.35 (2H, m), 10.85 (1H, brs), 13.57 (1H, brs)

Reference example 12-4

3-(4-Benzylpiperazine-1-sulfonyl)benzoic acid

¹H-NMR (DMSO-d₆) δ ppm: 2.35-2.50 (4H, m), 2.80-3.00 (4H, m),
3.46 (2H, s), 7.15-7.40 (5H, m), 7.75-7.85 (1H, m), 7.90-8.00
5 (1H, m), 8.15-8.30 (2H, m), 13.54 (1H, brs)

Reference example 13-1

Methyl 3-cyclohexyloxybenzoate

To a solution of cyclohexanol (1.10mL) in tetrahydrofuran
10 (10mL) was added triethylamine (1.50mL) under ice-cooling, and
then methansulfonyl chloride (0.81mL) was added to the reaction
mixture during 5 minutes. The resulting mixture was stirred at
room temperature for 30 minutes. Insoluble materials were removed
by filtration, and the filtrate was added to a mixture of methyl
15 3-hydroxybenzoate (1.52g), cesium carbonate (3.91g) and
N,N-dimethylformamide (5mL) under stirring. The mixture was
stirred at 60°C for 3 days. Water and diethyl ether were added
to the reaction mixture, and the organic layer was separated,
and washed with water, a saturated solution of sodium bicarbonate,
20 and brine successively. After the organic layer was dried over
anhydrous magnesium sulfate, the solvent was removed under
reduced pressure. The residue was purified by column
chromatography on silica gel (eluent: hexane/ethyl acetate =
3/1 - 1/1) to give the title compound (0.337g).

25 ¹H-NMR (CDCl₃) δ ppm: 1.20-1.65 (6H, m), 1.75-2.05 (4H, m), 3.91
(3H, s), 4.25-4.35 (1H, m), 7.05-7.15 (1H, m), 7.25-7.35 (1H,
m), 7.50-7.65 (2H, m)

Reference example 13-2

Reference example 13-2 was prepared in a manner similar to those as described in Reference Example 13-1 using

5 tetrahydropyran-4-ol instead of cyclohexanol.

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.75-1.85 (2H, m), 1.95-2.10 (2H, m), 3.55-3.65 (2H, m), 3.92 (3H, s), 3.95-4.05 (2H, m), 4.50-4.60 (1H, m), 7.05-7.15 (1H, m), 7.25-7.40 (1H, m), 7.55-7.70 (2H, m)

10

Reference example 14

3-(1-Benzylpiperidin-4-yloxy)benzoic acid

To a mixture of 1-benzylpiperidin-4-ol (2.29g), triphenylphosphine (3.14g) and tetrahydrofuran (20mL) were added diisopropyl azodicarboxylate (2.40mL) and methyl

15

3-hydroxybenzoate (1.52g) successively under ice-cooling, and the resulting mixture was stirred at room temperature under argon atmosphere overnight. Diethyl ether and 2mol/L hydrochloric acid were added to the reaction mixture. The aqueous layer was

20

separated and washed with diethyl ether. The aqueous layer was made alkaline with a saturated aqueous solution of sodium carbonate, and diethyl ether was added to the mixture. The organic layer was separated, washed with water and brine successively, and dried over anhydrous magnesium sulfate. The solvent was

25

removed under reduced pressure to give methyl 3-(1-benzylpiperidin-4-yloxy)benzoate. A 2mol/L aqueous solution of sodium hydroxide (20mL) was added to a mixture of

methyl 3-(1-benzylpiperidin-4-yloxy)benzoate and methanol (20mL), and the mixture was stirred at 60°C for 1 hour. After methanol was removed under reduced pressure, water and diethyl ether were added to the residue. The aqueous layer was separated, and washed with diethyl ether. Then, the aqueous layer was neutralized with 2mol/L hydrochloric acid, and the resulting precipitate was collected by filtration to give the title compound (2.56g).

¹H-NMR (CDCl₃) δ ppm: 1.90-2.20 (4H, m), 2.60-3.00 (4H, m), 3.79 (2H, s), 4.45-4.55 (1H, m), 7.00-7.10 (1H, m), 7.25-7.45 (6H, m), 7.55-7.70 (2H, m)

Reference example 15-1

Ethyl 6-(3-hydroxymethylphenoxy)nicotinate

A mixture of ethyl 6-chloronicotinate (1.0g), 3-hydroxymethylphenol (0.736g), potassium carbonate (1.49g) and 1-methyl-2-pyrrolidone (5mL) was stirred at 70°C for 2 days. Water and ethyl acetate were added to the reaction mixture. The organic layer was separated and washed with water, a saturated aqueous solution of sodium carbonate, water and brine successively. The organic layer was dried over anhydrous magnesium sulfate, and the solvent was removed under reduced pressure. The residue was purified by column chromatography on aminopropylated silica gel (eluent: hexane/ethyl acetate = 10/1 - 0/1) to give the title compound (0.44g).

¹H-NMR (CDCl₃) δ ppm: 1.38 (3H, t, J=7.2Hz), 1.83 (1H, t, J=6.0Hz), 4.38 (2H, q, J=7.2Hz), 4.73 (2H, d, J=6.0Hz), 6.90-7.00 (1H,

m), 7.05-7.10 (1H, m), 7.15-7.30 (2H, m), 7.35-7.45 (1H, m),
8.25-8.35 (1H, m), 8.75-8.85 (1H, m)

Reference examples 15-2 to 15-14 were prepared in a manner
5 similar to those as described in Reference Example 15-1 using
the corresponding phenols or pyridinols instead of
3-hydroxymethylphenol.

Reference example 15-2

Ethyl 6-(2-fluorophenoxy)nicotinate

10 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.38 (3H, t, $J=7.3\text{Hz}$), 4.37 (2H, q, $J=7.3\text{Hz}$),
7.00-7.10 (1H, m), 7.15-7.30 (4H, m), 8.25-8.35 (1H, m), 8.78
(1H, d, $J=1.8\text{Hz}$)

Reference example 15-3

Ethyl 6-(3-fluorophenoxy)nicotinate

15 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.39 (3H, t, $J=7.0\text{Hz}$), 4.39 (2H, q, $J=7.0\text{Hz}$),
6.85-7.05 (4H, m), 7.30-7.45 (1H, m), 8.25-8.35 (1H, m), 8.83
(1H, d, $J=2.3\text{Hz}$)

Reference example 15-4

Ethyl 6-(4-fluorophenoxy)nicotinate

20 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.38 (3H, t, $J=7.2\text{Hz}$), 4.38 (2H, q, $J=7.2\text{Hz}$),
6.90-7.00 (1H, m), 7.05-7.15 (4H, m), 8.25-8.35 (1H, m),
8.75-8.85 (1H, m)

Reference example 15-5

Ethyl 6-(2,4-difluorophenoxy)nicotinate

25 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.30-1.45 (3H, m), 4.30-4.45 (2H, m),
6.85-7.10 (3H, m), 7.10-7.25 (1H, m), 8.25-8.35 (1H, m), 8.77
(1H, d, $J=2.3\text{Hz}$)

Reference example 15-6

Ethyl 6-(2,4,6-trifluorophenoxy)nicotinate

¹H-NMR (CDCl₃) δ ppm: 1.38 (3H, t, J=7.2Hz), 4.38 (2H, q, J=7.2Hz),
6.75-6.85 (2H, m), 7.05-7.20 (1H, m), 8.30-8.40 (1H, m), 8.75
5 (1H, d, J=2.2Hz)

Reference example 15-7

Ethyl 6-(2-isopropylphenoxy)nicotinate

¹H-NMR (CDCl₃) δ ppm: 1.18 (6H, d, J=6.9Hz), 1.38 (3H, t, J=7.2Hz),
3.00-3.15 (1H, m), 4.37 (2H, q, J=7.2Hz), 6.85-6.95 (1H, m),
10 7.00-7.10 (1H, m), 7.20-7.30 (2H, m), 7.35-7.45 (1H, m),
8.20-8.30 (1H, m), 8.80-8.90 (1H, m)

Reference example 15-8

Ethyl 6-(4-isopropylphenoxy)nicotinate

¹H-NMR (CDCl₃) δ ppm: 1.27 (6H, d, J=6.9Hz), 1.38 (3H, t, J=7.3Hz),
15 2.85-3.00 (1H, m), 4.30-4.45 (2H, m), 6.91 (1H, d, J=8.5Hz),
7.00-7.15 (2H, m), 7.20-7.35 (2H, m), 8.20-8.30 (1H, m),
8.80-8.90 (1H, m)

Reference example 15-9

Ethyl 6-(4-fluoro-2-methylphenoxy)nicotinate

¹H-NMR (CDCl₃) δ ppm: 1.38 (3H, t, J=7.1Hz), 2.14 (3H, s), 4.30-4.45
20 (2H, m), 6.85-7.10 (4H, m), 8.25-8.35 (1H, m), 8.75-8.85 (1H,
m),

Reference example 15-10

Ethyl 6-(2-methoxyphenoxy)nicotinate

¹H-NMR (CDCl₃) δ ppm: 1.37 (3H, t, J=7.2Hz), 3.75 (3H, s), 4.36
25 (2H, q, J=7.2Hz), 6.90-6.97 (1H, m), 6.98-7.10 (2H, m), 7.10-7.20
(1H, m), 7.20-7.30 (1H, m), 8.20-8.30 (1H, m), 8.79 (1H, d,

J=2.7Hz)

Reference example 15-11

Ethyl 6-(3-methoxyphenoxy)nicotinate

¹H-NMR (CDCl₃) δ ppm: 1.38 (3H, t, J=7.1Hz), 3.81 (3H, s), 4.38
5 (2H, q, J=7.1Hz), 6.65-6.85 (3H, m), 6.92 (1H, d, J=8.7Hz), 7.32
(1H, t, J=8.2Hz), 8.27 (1H, dd, J=2.3, 8.7Hz), 8.84 (1H, d,
J=2.3Hz)

Reference example 15-12

Ethyl 6-(4-fluoro-2-methoxyphenoxy)nicotinate

10 ¹H-NMR (CDCl₃) δ ppm: 1.37 (3H, t, J=7.2Hz), 3.74 (3H, s), 4.37
(2H, m), 6.65-6.80 (2H, m), 6.90-7.00 (1H, m), 7.05-7.15 (1H,
m), 8.20-8.30 (1H, m), 8.77 (1H, d, J=2.1Hz)

Reference example 15-13

Ethyl 6-(pyridin-3-yloxy)nicotinate

15 ¹H-NMR (CDCl₃) δ ppm: 1.39 (3H, t, J=7.2Hz), 4.39 (2H, q, J=7.2Hz),
7.00-7.10 (1H, m), 7.35-7.45 (1H, m), 7.50-7.60 (1H, m),
8.30-8.40 (1H, m), 8.45-8.60 (2H, m), 8.75-8.85 (1H, m)

Reference example 15-14

Ethyl 6-phenethyloxynicotinate

20 ¹H-NMR (CDCl₃) δ ppm: 1.35-1.45 (3H, m), 3.00-3.15 (2H, m),
4.30-4.65 (4H, m), 6.65-6.80 (1H, m), 7.15-7.40 (5H, m),
8.05-8.20 (1H, m), 8.75-8.90 (1H, m)

Reference example 16

25 Methyl 4-hydroxy-3-morpholin-4-ylmethylbenzoate

Concentrated sulfuric acid (1.2mL) was added to a solution
of 4-hydroxy-3-morpholin-4-ylmethylbenzoic acid mono hydrate

(5.11g) in methanol (50mL), and the resulting mixture was heated under reflux overnight. A saturated aqueous solution of sodium carbonate and ethyl acetate were added to the reaction mixture. The organic layer was separated, washed with brine, and dried
 5 over anhydrous magnesium sulfate. The solvent was removed under reduced pressure to give the title compound (3.86g).

¹H-NMR (CDCl₃) δ ppm: 2.40-2.80 (4H, m), 3.60-3.85 (6H, m), 3.87 (3H, s), 6.80-6.90 (1H, m), 7.70-7.80 (1H, m), 7.85-7.95 (1H, m)

10

Reference example 17-1

Methyl 4-benzyloxy-3-morpholin-4-ylmethylbenzoate

To a mixture of methyl 4-hydroxy-3-morpholin-4-ylmethylbenzoate (3.85g), potassium carbonate (4.24g) and
 15 N,N-dimethylformamide (15mL) was added benzyl bromide (1.85mL), and the resulting mixture was stirred at room temperature for 3 days. Water was added to the reaction mixture, and the resulting precipitate was collected by filtration to give the title compound (4.69g).

20 ¹H-NMR (CDCl₃) δ ppm: 2.40-2.60 (4H, m), 3.61 (2H, brs), 3.65-3.80 (4H, m), 3.89 (3H, s), 5.15 (2H, s), 6.90-7.00 (1H, m), 7.30-7.50 (5H, m), 7.90-8.10 (2H, m)

Reference Example 17-2

25 Ethyl 4-ethoxy-3-morpholin-4-ylmethylbenzoate

Reference Example 17-2 was prepared in a manner similar to those as described in Reference Example 17-1 using

4-hydroxy-3-morpholin-4-ylmethylbenzoic acid mono hydrate and ethyl iodide instead of methyl 4-hydroxy-3-morpholin-4-ylmethylbenzoate and benzyl bromide.

¹H-NMR (CDCl₃) δ ppm: 1.39 (3H, t, J=7.1Hz), 1.45 (3H, t, J=7.0Hz),
 5 2.45-2.60 (4H, m), 3.57 (2H, s), 3.65-3.80 (4H, m), 4.10 (2H, q, J=7.0Hz), 4.35 (2H, q, J=7.1Hz), 6.80-6.90 (1H, m), 7.90-8.10 (2H, m)

Reference Example 18-1

10 6-(3-Chloromethylphenoxy)nicotinoyl chloride hydrochloride

To a mixture of 6-(3-hydroxymethylphenoxy)nicotinic acid (0.4g) and toluene (10mL) were added thionyl chloride (0.715mL) and N,N-dimethylformamide (0.040mL) successively, and the resulting mixture was stirred at 80°C for 2 hours. The reaction
 15 mixture was concentrated under reduced pressure to give the title compound (0.42g).

¹H-NMR (DMSO-d₆) δ ppm: 4.63 (2H, s), 6.95-7.00 (1H, m), 7.05-7.45 (4H, m), 7.62 (1H, s), 8.25-8.35 (1H, m), 8.75-8.80 (1H, m)

20 Reference Examples 18-2 to 18-19 were prepared in a manner similar to those as described in Reference Example 18-1 using the corresponding carboxylic acids instead of 6-(3-hydroxymethylphenoxy)nicotinic acid.

Reference Example 18-2

25 6-(2-Fluorophenoxy)nicotinoyl chloride

¹H-NMR (CDCl₃) δ ppm: 7.11 (1H, d, J=8.8Hz), 7.15-7.35 (4H, m), 8.30-8.40 (1H, m), 8.88 (1H, d, J=2.0Hz)

Reference example 18-3

6-(3-Fluorophenoxy)nicotinoyl chloride

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 6.85-7.10 (4H, m), 7.35-7.45 (1H, m),
8.30-8.40 (1H, m), 8.92 (1H, d, $J=2.6\text{Hz}$)

5 Reference example 18-4

6-(4-Fluorophenoxy)nicotinoyl chloride

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 7.03 (1H, d, $J=8.6\text{Hz}$), 7.10-7.20 (4H, m),
8.30-8.40 (1H, m), 8.90 (1H, d, $J=2.1\text{Hz}$)

Reference example 18-5

10 6-(2,4-Difluorophenoxy)nicotinoyl chloride

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 6.90-7.05 (2H, m), 7.10-7.15 (1H, m),
7.15-7.25 (1H, m), 8.30-8.40 (1H, m), 8.85-8.90 (1H, m)

Reference example 18-6

6-(2,4-Trifluorophenoxy)nicotinoyl chloride

15 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 6.75-6.90 (2H, m), 7.20 (1H, d, $J=8.6\text{Hz}$),
8.35-8.45 (1H, m), 8.85 (1H, d, $J=2.2\text{Hz}$)

Reference example 18-7

6-(2-Isopropyl-phenoxy)nicotinoyl chloride

20 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.19 (6H, m), 2.95-3.10 (1H, m), 6.95-7.10
(2H, m), 7.20-7.30 (2H, m), 7.35-7.45 (1H, m), 8.25-8.40 (1H,
m), 8.91 (1H, d, $J=2.5\text{Hz}$)

Reference example 18-8

6-(4-Isopropyl-phenoxy)nicotinoyl chloride

25 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.28 (6H, d, $J=6.9\text{Hz}$), 2.90-3.05 (1H, m),
6.99 (1H, d, $J=8.9\text{Hz}$), 7.00-7.15 (2H, m), 7.25-7.35 (2H, m),
8.25-8.35 (1H, m), 8.94 (1H, d, $J=2.5\text{Hz}$)

Reference example 18-9

6-(4-Fluoro-2-methylphenoxy)nicotinoyl chloride

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 2.14 (3H, s), 6.90-7.10 (4H, m), 8.30-8.40 (1H, m), 8.89 (1H, d, $J=2.4\text{Hz}$)

Reference example 18-10

5 6-(2-Methoxyphenoxy)nicotinoyl chloride

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 3.76 (3H, s), 6.95-7.10 (3H, m), 7.10-7.20 (1H, m), 7.20-7.35 (1H, m), 8.25-8.35 (1H, m), 8.85-8.95 (1H, m)

Reference example 18-11

10 6-(3-Methoxyphenoxy)nicotinoyl chloride

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 3.82 (3H, s), 6.65-6.80 (2H, m), 6.80-6.90 (1H, m), 7.00 (1H, d, $J=8.5\text{Hz}$), 7.35 (1H, t, $J=8.2\text{Hz}$), 8.25-8.40 (1H, m), 8.93 (1H, d, $J=2.5\text{Hz}$)

Reference example 18-12

15 6-(4-Fluoro-2-methoxyphenoxy)nicotinoyl chloride

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 3.74 (3H, s), 6.60-6.80 (2H, m), 7.00-7.15 (2H, m), 8.25-8.35 (1H, m), 8.85-8.90 (2H, m)

Reference example 18-13

5-(3-Chloromethylphenyl)furan-2-carbonyl chloride

20 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 7.18 (1H, d, $J=3.5\text{Hz}$), 7.33 (1H, d, $J=3.5\text{Hz}$), 7.40-7.55 (2H, m), 7.75-7.85 (1H, m), 7.85-7.95 (1H, m)

Reference example 18-14

5-Pyridin-2-ylfuran-2-carbonyl chloride

25 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 7.60-7.70 (1H, m), 7.85-7.95 (1H, m), 8.35-8.55 (2H, m), 8.75-8.90 (2H, m)

Reference example 18-15

5-Pyridin-3-ylfuran-2-carbonyl chloride

¹H-NMR (CDCl₃) δ ppm: 7.35-7.50 (2H, m), 7.75-7.85 (1H, m),
8.45-8.55 (1H, m), 8.65-8.75 (1H, m), 9.10-9.20 (1H, m)

Reference example 18-16

5-Pyridin-4-ylfuran-2-carbonyl chloride

5 ¹H-NMR (CDCl₃) δ ppm: 7.45-7.55 (1H, m), 7.85-7.90 (1H, m),
8.25-8.35 (2H, m), 8.85-8.95 (2H, m)

Reference example 18-17

5-(Azepane-1-sulfonyl)furan-2-carbonyl chloride

¹H-NMR (CDCl₃) δ ppm: 1.55-1.85 (8H, m), 3.35-3.50 (4H, m),
10 7.00-7.10 (1H, m), 7.45-7.50 (1H, m)

Reference example 18-18

3-(4-Methylpiperazine-1-sulfonyl)benzoyl chloride
hydrochloride

¹H-NMR (DMSO-d₆) δ ppm: 2.60-2.85 (5H, m), 3.05-3.25 (2H, m),
15 3.35-3.50 (2H, m), 3.70-3.90 (2H, m), 7.80-7.90 (1H, m),
8.00-8.10 (1H, m), 8.20-8.35 (2H, m), 10.55 (1H, brs)

Reference example 18-19

4-Benzyloxy-3-morpholin-4-ylmethylbenzoyl chloride

¹H-NMR (DMSO-d₆) δ ppm: 2.80-2.95 (2H, m), 3.25-3.40 (2H, m),
20 3.80-3.95 (2H, m), 4.20-4.40 (2H, m), 4.31 (2H, s), 5.27 (2H,
brs), 7.10-7.20 (1H, m), 7.35-7.50 (5H, m), 8.15-8.25 (1H, m),
8.45-8.50 (1H, m), 13.38 (1H, brs)

Reference example 19-1

25 5-Pyridin-2-ylfuran-2-carboxylic acid

To a solution of silver nitrate (0.888g) in water (5mL)
was added a 2mol/L aqueous solution of sodium hydroxide (5mL)

under ice-cooling. Then, 5-pyridin-2-ylfuran-2-carbaldehyde (0.376g) and tetrahydrofuran (1mL) were added to the mixture, and the resulting mixture was stirred at room temperature for 1 hour. Insoluble materials were removed by filtration. To the
 5 filtrate was added 2mol/L hydrochloric acid (5mL) under ice-cooling, and the mixture was concentrated under reduced pressure. The resulting precipitate was collected by filtration to give the title compound (0.289g).

¹H-NMR (DMSO-d₆) δ ppm: 7.25-7.50 (3H, m), 7.85-8.05 (2H, m),
 10 8.60-8.70 (1H, m)

Reference Examples 19-2 and 19-3 were prepared in a manner similar to those as described in Reference Example 19-1 using the corresponding aldehydes instead of 5-pyridin-2-ylfuran-
 15 2-carbaldehyde.

Reference example 19-2

5-(3-Hydroxymethylphenyl)furan-2-carboxylic acid

¹H-NMR (DMSO-d₆) δ ppm: 4.50-4.60 (2H, m), 5.20-5.40 (1H, m),
 7.12 (1H, d, J=3.9Hz), 7.25-7.40 (2H, m), 7.43 (1H, t, J=7.8Hz),
 20 7.68 (1H, d, J=7.8Hz), 7.77 (1H, s), 13.11 (1H, brs)

Reference example 19-3

5-Cyclopropylfuran-2-carboxylic acid

¹H-NMR (CDCl₃) δ ppm: 0.85-1.05 (4H, m), 1.95-2.05 (1H, m), 6.10
 (1H, d, J=3.5Hz), 7.22 (1H, d, J=3.5Hz)

25

Example 1

3-Cyclopropanecarbonylamino-5-fluorobenzofuran-2-

carboxamide (compound 1-1)

3-Amino-5-fluorobenzofuran-2-carboxamide (0.1g) was dissolved in tetrahydrofuran (3mL), and the solution was cooled at 0°C. Then, triethylamine (0.144mL) and cyclopropanecarbonyl chloride (0.051mL) was added to the mixture successively, and the resulting mixture was stirred at room temperature for 18 hours. Water (6mL) was added to the reaction mixture, and the mixture was stirred for another 1 hour. The resulting precipitate was collected by filtration to give the title compound (0.063g).

¹H-NMR (DMSO-d₆) δ ppm: 0.80-0.98 (4H, m), 1.91-2.06 (1H, m), 7.28-7.40 (1H, m), 7.55-7.70 (2H, m), 7.89 (1H, brs), 8.13 (1H, brs), 10.34 (1H, brs)

Example 2

3-Cyclopropanecarbonylamino-5-chlorobenzofuran-2-carboxamide (compound 1-2)

Cyclopropanecarboxylic acid (0.018g) and triphosgen (0.032g) were dissolved in tetrahydrofuran (1.5mL), and N-methylmorpholine (0.047mL) was added thereto at room temperature. The mixture was stirred for 15 minutes. Then, a solution of 3-amino-5-chlorobenzofuran-2-carboxamide (0.042g) in tetrahydrofuran (1.5mL) was added to the mixture, and the resulting mixture was stirred for 14 hours. The reaction mixture was extracted with diethyl ether, washed with a 10% aqueous solution of sodium hydroxide and brine successively. The organic solvent was removed under reduced pressure, and the residue was purified by column chromatography on silica gel (eluent:

hexane/tetrahydrofuran = 7/3) to give the title compound (0.013g).

¹H-NMR (DMSO-d₆) δ ppm: 0.80-0.95 (4H, m), 1.95-2.05 (1H, m),
7.45-7.65 (2H, m), 7.92 (1H, brs), 7.95-8.00 (1H, m), 8.15 (1H,
5 brs), 10.37 (1H, brs)

Compounds 1-3 to 1-370 were prepared in a manner similar to those as described in Example 1 or Example 2 using the corresponding 3-aminobenzofuran-2-carboxamides and the
10 corresponding acid chlorides or carboxylic acids instead of 3-amino-5-fluorobenzofuran-2-carboxamide and cyclopropanecarbonylchloride. These were illustrated in table 2.

Table 2

Compound No.	Structure	Compound No.	Structure
1-1		1-9	
1-2		1-10	
1-3		1-11	
1-4		1-12	
1-5		1-13	
1-6		1-14	
1-7		1-15	
1-8		1-16	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-17		1-25	
1-18		1-26	
1-19		1-27	
1-20		1-28	
1-21		1-29	
1-22		1-30	
1-23		1-31	
1-24		1-32	

Table 2 (continued)

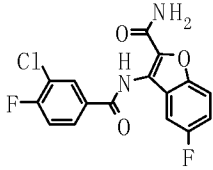
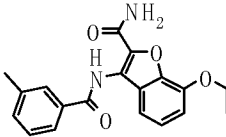
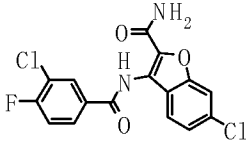
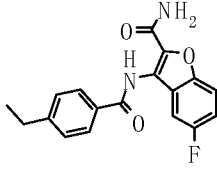
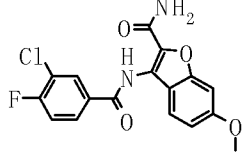
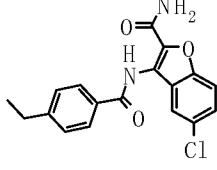
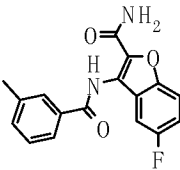
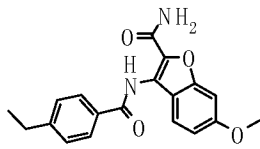
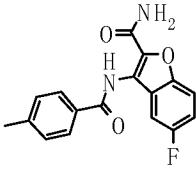
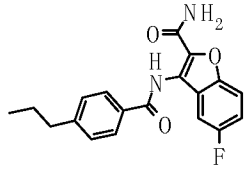
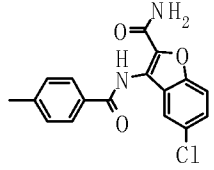
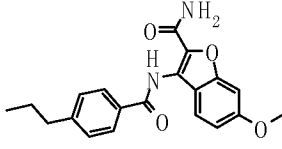
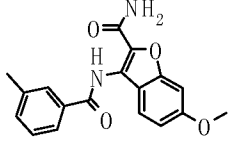
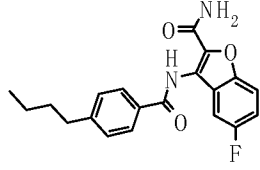
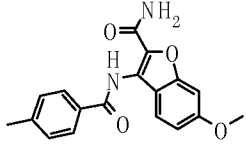
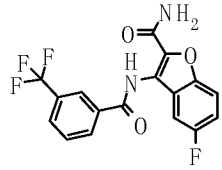
Compound No.	Structure	Compound No.	Structure
1-33		1-41	
1-34		1-42	
1-35		1-43	
1-36		1-44	
1-37		1-45	
1-38		1-46	
1-39		1-47	
1-40		1-48	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-49		1-57	
1-50		1-58	
1-51		1-59	
1-52		1-60	
1-53		1-61	
1-54		1-62	
1-55		1-63	
1-56		1-64	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-65		1-73	
1-66		1-74	
1-67		1-75	
1-68		1-76	
1-69		1-77	
1-70		1-78	
1-71		1-79	
1-72		1-80	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-81		1-89	
1-82		1-90	
1-83		1-91	
1-84		1-92	
1-85		1-93	
1-86		1-94	
1-87		1-95	
1-88		1-96	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-97		1-105	
1-98		1-106	
1-99		1-107	
1-100		1-108	
1-101		1-109	
1-102		1-110	
1-103		1-111	
1-104		1-112	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-113		1-121	
1-114		1-122	
1-115		1-123	
1-116		1-124	
1-117		1-125	
1-118		1-126	
1-119		1-127	
1-120		1-128	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-129		1-137	
1-130		1-138	
1-131		1-139	
1-132		1-140	
1-133		1-141	
1-134		1-142	
1-135		1-143	
1-136		1-144	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-145		1-154	
1-146		1-155	
1-147		1-156	
1-148		1-157	
1-149		1-158	
1-150		1-159	
1-151		1-160	
1-152		1-161	
1-153		1-162	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-163		1-171	
1-164		1-172	
1-165		1-173	
1-166		1-174	
1-167		1-175	
1-168		1-176	
1-169		1-177	
1-170		1-178	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-179		1-187	
1-180		1-188	
1-181		1-189	
1-182		1-190	
1-183		1-191	
1-184		1-192	
1-185		1-193	
1-186		1-194	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-195		1-203	
1-196		1-204	
1-197		1-205	
1-198		1-206	
1-199		1-207	
1-200		1-208	
1-201		1-209	
1-202		1-210	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-211		1-219	
1-212		1-220	
1-213		1-221	
1-214		1-222	
1-215		1-223	
1-216		1-224	
1-217		1-225	
1-218		1-226	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-227		1-235	
1-228		1-236	
1-229		1-237	
1-230		1-238	
1-231		1-239	
1-232		1-240	
1-233		1-241	
1-234		1-242	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-243		1-251	
1-244		1-252	
1-245		1-253	
1-246		1-254	
1-247		1-255	
1-248		1-256	
1-249		1-257	
1-250		1-258	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-259		1-267	
1-260		1-268	
1-261		1-269	
1-262		1-270	
1-263		1-271	
1-264		1-272	
1-265		1-273	
1-266		1-274	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-275		1-283	
1-276		1-284	
1-277		1-285	
1-278		1-286	
1-279		1-287	
1-280		1-288	
1-281		1-289	
1-282		1-290	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-291		1-299	
1-292		1-300	
1-293		1-301	
1-294		1-302	
1-295		1-303	
1-296		1-304	
1-297		1-305	
1-298		1-306	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-307		1-315	
1-308		1-316	
1-309		1-317	
1-310		1-318	
1-311		1-319	
1-312		1-320	
1-313		1-321	
1-314		1-322	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-323		1-331	
1-324		1-332	
1-325		1-333	
1-326		1-334	
1-327		1-335	
1-328		1-336	
1-329		1-337	
1-330		1-338	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-339		1-347	
1-340		1-348	
1-341		1-349	
1-342		1-350	
1-343		1-351	
1-344		1-352	
1-345		1-353	
1-346		1-354	

Table 2 (continued)

Compound No.	Structure	Compound No.	Structure
1-355		1-363	
1-356		1-364	
1-357		1-365	
1-358		1-366	
1-359		1-367	
1-360		1-368	
1-361		1-369	
1-362		1-370	

The physical data of compounds 1-3 to 1-370 were described below.

Compound 1-3

¹H-NMR (CDCl₃) δ ppm: 1.65-2.45 (4H, m), 3.95-4.10 (1H, m),
5 4.15-4.25 (1H, m), 4.50-4.65 (1H, m), 6.47 (1H, brs), 6.70 (1H, brs), 7.30-7.50 (2H, m), 8.55-8.65 (1H, m), 10.89 (1H, brs)

Compound 1-4

¹H-NMR (DMSO-d₆) δ ppm: 3.87 (3H, s), 7.00 (1H, dd, J=2.2, 8.8Hz),
7.10 (1H, d, J=2.2Hz), 7.55-7.75 (3H, m), 7.76-8.20 (4H, m),
10 8.26 (1H, d, J=8.8Hz), 11.18 (1H, s)

Compound 1-5

¹H-NMR (DMSO-d₆) δ ppm: 7.36-7.95 (7H, m), 8.01 (1H, brs), 8.28 (1H, brs), 10.98 (1H, s)

Compound 1-6

15 ¹H-NMR (DMSO-d₆) δ ppm: 7.20-7.35 (1H, m), 7.40-7.50 (2H, m), 7.55 (1H, dd, J=2.2, 9.1Hz), 7.95 (1H, brs), 8.00-8.13 (2H, m), 8.15-8.30 (2H, m), 11.03 (1H, s)

Compound 1-7

¹H-NMR (DMSO-d₆) δ ppm: 7.38-7.90 (6H, m), 7.99 (1H, brs),
20 8.15-8.32 (2H, m), 11.03 (1H, s)

Compound 1-8

¹H-NMR (MeOD-d₄) δ ppm: 7.20-7.35 (2H, m), 7.40-7.55 (2H, m), 8.05-8.15 (2H, m), 8.45-8.55 (1H, m)

Compound 1-9

25 ¹H-NMR (DMSO-d₆) δ ppm: 7.45-7.95 (6H, m), 8.06 (1H, brs), 8.33 (1H, brs), 10.94 (1H, s)

Compound 1-10

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.48–7.94 (6H, m), 8.07 (1H, brs), 8.18 (1H, brs), 10.91 (1H, s)

Compound 1-11

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.43 (3H, s), 7.30–7.58 (3H, m), 7.62–7.96 (5H, m), 8.19 (1H, brs), 10.95 (1H, s)

Compound 1-12

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.54 (3H, s), 7.20–7.38 (2H, m), 7.48–7.72 (2H, m), 7.76–8.00 (4H, m), 8.17 (1H, brs), 10.94 (1H, s)

Compound 1-13

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.50–7.93 (4H, m), 8.10–8.50 (5H, m), 10.96 (1H, s)

Compound 1-14

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.81 (3H, s), 7.09–7.19 (1H, m), 7.45–7.72 (4H, m), 7.77–8.00 (3H, m), 8.18 (1H, brs), 10.94 (1H, s)

Compound 1-15

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.99 (3H, s), 7.08–7.32 (2H, m), 7.48–7.72 (3H, m), 7.76–8.00 (3H, m), 8.13 (1H, brs), 10.93 (1H, s)

Compound 1-16

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.99 (3H, s), 7.13 (1H, d, $J=7.9\text{Hz}$), 7.20–7.30 (1H, m), 7.35–7.50 (2H, m), 7.70 (1H, d, $J=7.9\text{Hz}$), 7.89 (1H, brs), 8.00–8.20 (3H, m), 10.91 (1H, s)

Compound 1-17

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.87 (3H, s), 6.92–7.20 (2H, m), 7.45–8.28 (7H, m), 11.14 (1H, s)

Compound 1-18

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.43 (3H, t, $J=6.9\text{Hz}$), 4.30 (2H, q, $J=6.9\text{Hz}$), 7.05–7.15 (1H, m), 7.20–7.30 (1H, m), 7.35–7.50 (2H,

m), 7.66 (1H, dd, J=0.9, 8.2Hz), 7.90 (1H, brs), 8.00-8.20 (3H, m), 10.89 (1H, s)

Compound 1-19

¹H-NMR (DMSO-d₆) δ ppm: 5.24 (2H, s), 7.00-7.20 (2H, m), 7.30-7.95
5 (10H, m), 8.11 (1H, brs), 8.17 (1H, d, J=9.0Hz), 11.14 (1H, s)

Compound 1-20

¹H-NMR (DMSO-d₆) δ ppm: 7.49-8.24 (10H, m), 8.37 (1H, brs), 8.50
(1H, d, J=8.2Hz), 11.03 (1H, s)

Compound 1-21

10 ¹H-NMR (DMSO-d₆) δ ppm: 1.14 (6H, t, J=7.1Hz), 3.43 (4H, q, J=7.1Hz), 6.58-6.64 (1H, m), 6.76-6.86 (1H, m), 7.48-7.92 (6H, m), 8.11 (1H, d, J=9.2Hz), 11.27 (1H, s)

Compound 1-22

¹H-NMR (CDCl₃) δ ppm: 5.64 (1H, brs), 6.43 (1H, brs), 7.20-7.65
15 (4H, m), 7.85-7.95 (1H, m), 8.00-8.10 (1H, m), 8.30-8.40 (1H, m), 10.88 (1H, brs)

Compound 1-23

¹H-NMR (CDCl₃) δ ppm: 5.99 (1H, brs), 6.52 (1H, brs), 7.15-7.30
20 (1H, m), 7.35-7.55 (3H, m), 7.95-8.05 (2H, m), 8.30-8.40 (1H, m), 10.93 (1H, brs)

Compound 1-24

¹H-NMR (DMSO-d₆) δ ppm: 7.57 (1H, dd, J=2.2, 8.8Hz), 7.60-7.70
(2H, m), 7.72-7.80 (1H, m), 7.90-8.10 (3H, m), 8.19 (1H, d, J=2.2Hz), 8.27 (1H, brs), 10.99 (1H, s)

25 Compound 1-25

¹H-NMR (DMSO-d₆) δ ppm: 7.36-7.48 (1H, m), 7.64-7.80 (3H, m),
7.92-8.08 (3H, m), 8.16-8.32 (2H, m), 11.04 (1H, s)

Compound 1-26

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.52-7.80 (4H, m), 7.96-8.12 (3H, m),
8.33 (1H, brs), 10.95 (1H, s)

Compound 1-27

5 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.87 (3H, s), 6.95-7.05 (1H, m), 7.07-7.15
(1H, m), 7.55-8.25 (7H, m), 11.13 (1H, s)

Compound 1-28

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.87 (3H, s), 7.00 (1H, dd, $J=2.2, 8.8\text{Hz}$),
7.11 (1H, d, $J=2.2\text{Hz}$), 7.65-7.75 (2H, m), 7.76-8.25 (5H, m),
10 11.15 (1H, s)

Compound 1-29

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.35-7.45 (1H, m), 7.50-7.75 (2H, m),
7.80-8.05 (4H, m), 8.15-8.30 (2H, m), 10.96 (1H, s)

Compound 1-30

15 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.87 (3H, s), 7.00 (1H, dd, $J=2.2, 8.8\text{Hz}$),
7.11 (1H, d, $J=2.2\text{Hz}$), 7.50-7.65 (1H, m), 7.75-8.25 (6H, m),
11.12 (1H, s)

Compound 1-31

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.35-7.45 (1H, m), 7.60-7.75 (2H, m),
20 7.80-8.10 (4H, m), 8.24 (1H, brs), 10.91 (1H, s)

Compound 1-32

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.87 (3H, s), 6.99 (1H, dd, $J=2.2, 8.8\text{Hz}$),
7.11 (1H, d, $J=2.2\text{Hz}$), 7.60-8.20 (6H, m), 11.06 (1H, s)

Compound 1-33

25 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.35-7.45 (1H, m), 7.60-7.70 (2H, m),
7.75-7.85 (1H, m), 7.96 (1H, brs), 8.00-8.10 (1H, m), 8.15-8.30
(2H, m), 10.90 (1H, brs)

Compound 1-34

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.40–7.45 (1H, m), 7.60–7.70 (1H, m), 7.75–7.80 (1H, m), 7.97 (1H, brs), 8.00–8.30 (4H, m), 10.95 (1H, brs)

5 Compound 1-35

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.87 (3H, s), 6.95–7.05 (1H, m), 7.10–7.15 (1H, m), 7.60–7.70 (1H, m), 7.85 (1H, brs), 7.95–8.25 (4H, m), 11.06 (1H, brs)

Compound 1-36

10 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.43 (3H, s), 7.35–7.55 (3H, m), 7.60–7.70 (1H, m), 7.75–7.90 (2H, m), 7.95–8.10 (2H, m), 8.27 (1H, brs), 11.00 (1H, s)

Compound 1-37

15 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.42 (3H, s), 7.35–7.45 (3H, m), 7.66 (1H, m), 7.91 (2H, d, $J=8.2\text{Hz}$), 7.95–8.10 (2H, m), 8.27 (1H, brs), 11.01 (1H, s)

Compound 1-38

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.42 (3H, s), 7.35–7.70 (4H, m), 7.85–8.00 (2H, m), 8.03 (1H, brs), 8.30–8.40 (1H, m), 8.32 (1H, brs), 11.04
20 (1H, brs)

Compound 1-39

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.42 (3H, s), 3.87 (3H, s), 7.00 (1H, dd, $J=2.2, 9.1\text{Hz}$), 7.10 (1H, d, $J=2.2\text{Hz}$), 7.45–7.55 (2H, m), 7.74–8.20 (4H, m), 8.25 (1H, d, $J=8.8\text{Hz}$), 11.15 (1H, s)

25 Compound 1-40

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.41 (3H, s), 3.87 (3H, s), 6.99 (1H, dd, $J=2.5, 9.1\text{Hz}$), 7.10 (1H, d, $J=2.2\text{Hz}$), 7.41 (2H, d, $J=7.9\text{Hz}$),

7.70-8.20 (4H, m), 8.27 (1H, d, J=9.1Hz), 11.15 (1H, s)

Compound 1-41

¹H-NMR (DMSO-d₆) δ ppm: 1.43 (3H, t, J=6.9Hz), 2.42 (3H, s),
4.29 (2H, d, J=7.3Hz), 7.10-7.15 (1H, m), 7.20-7.30 (1H, m),
5 7.40-7.55 (2H, m), 7.73 (1H, dd, J=0.9, 8.2Hz), 7.76-7.85 (2H,
m), 7.90 (1H, brs), 8.08 (1H, brs), 10.93 (1H, s)

Compound 1-42

¹H-NMR (DMSO-d₆) δ ppm: 1.23 (3H, t, J=7.6Hz), 2.72 (2H, q,
J=7.6Hz), 7.35-7.50 (3H, m), 7.66 (1H, m), 7.85-8.10 (4H, m),
10 8.27 (1H, brs), 11.01 (1H, s)

Compound 1-43

¹H-NMR (DMSO-d₆) δ ppm: 1.23 (3H, t, J=7.6Hz), 2.72 (2H, q,
J=7.6Hz), 7.45 (2H, d, J=8.2Hz), 7.57 (1H, dd, J=2.5, 9.1Hz),
7.67 (1H, d, J=8.8Hz), 7.93 (2H, d, J=8.2Hz), 8.02 (1H, brs),
15 8.29 (1H, brs), 8.35 (1H, d, J=2.2Hz), 11.03 (1H, s)

Compound 1-44

¹H-NMR (DMSO-d₆) δ ppm: 1.23 (3H, t, J=7.6Hz), 2.71 (2H, q,
J=7.6Hz), 3.87 (3H, s), 6.99 (1H, dd, J=2.2, 9.1Hz), 7.10 (1H,
d, J=2.2Hz), 7.44 (2H, d, J=8.2Hz), 7.70-8.20 (4H, m), 8.27 (1H,
20 d, J=8.8Hz), 11.15 (1H, s)

Compound 1-45

¹H-NMR (DMSO-d₆) δ ppm: 0.92 (3H, t, J=7.3Hz), 1.55-1.75 (2H,
m), 2.66 (2H, t, J=7.3Hz), 7.35-7.50 (3H, m), 7.66 (1H, dd, J=4.1,
9.1Hz), 7.85-8.10 (4H, m), 8.27 (1H, brs), 11.00 (1H, s)

25 Compound 1-46

¹H-NMR (DMSO-d₆) δ ppm: 0.92 (3H, t, J=7.3Hz), 1.55-1.70 (2H,
m), 2.66 (2H, t, J=7.3Hz), 3.87 (3H, s), 6.99 (1H, dd, J=2.2,

8.8Hz), 7.10 (1H, d, J=2.2Hz), 7.42 (2H, d, J=8.2Hz), 7.70-8.20
(4H, m), 8.27 (1H, d, J=8.8Hz), 11.14 (1H, s)

Compound 1-47

¹H-NMR (DMSO-d₆) δ ppm: 0.91 (3H, t, J=7.3Hz), 1.25-1.40 (2H,
5 m), 1.55-1.65 (2H, m), 2.69 (2H, t, J=7.6Hz), 7.35-7.50 (3H,
m), 7.60-7.70 (1H, m), 7.85-8.10 (4H, m), 8.26 (1H, brs), 11.00
(1H, s)

Compound 1-48

¹H-NMR (DMSO-d₆) δ ppm: 7.35-7.45 (1H, m), 7.60-7.75 (1H, m),
10 7.80-7.90 (2H, m), 7.95-8.10 (2H, m), 8.20-8.40 (3H, m), 11.07
(1H, s)

Compound 1-49

¹H-NMR (DMSO-d₆) δ ppm: 3.87 (3H, s), 7.00 (1H, dd, J=2.2, 8.8Hz),
7.12 (1H, d, J=2.2Hz), 7.70-8.20 (5H, m), 8.25-8.35 (2H, m),
15 11.23 (1H, s)

Compound 1-50

¹H-NMR (DMSO-d₆) δ ppm: 4.87 (2H, s), 7.30-7.50 (1H, m), 7.60-7.75
(3H, m), 7.90-8.10 (4H, m), 8.27 (1H, brs), 11.02 (1H, s)

Compound 1-51

20 ¹H-NMR (CDCl₃) δ ppm: 4.68 (2H, s), 5.61 (1H, brs), 6.41 (1H,
brs), 7.20-7.30 (1H, m), 7.35-7.45 (1H, m), 7.50-7.70 (2H, m),
7.95-8.10 (2H, m), 8.35-8.45 (1H, m), 10.89 (1H, brs)

Compound 1-52

25 ¹H-NMR (DMSO-d₆) δ ppm: 4.87 (2H, s), 7.40-7.50 (1H, m), 7.60-7.80
(3H, m), 7.95-8.05 (3H, m), 8.20-8.30 (2H, m), 11.05 (1H, brs)

Compound 1-53

¹H-NMR (DMSO-d₆) δ ppm: 4.89 (2H, s), 7.40-7.50 (1H, m), 7.60-7.65

(1H, m), 7.70-7.80 (2H, m), 7.95-8.15 (3H, m), 8.20-8.30 (2H, m), 11.06 (1H, brs)

Compound 1-54

¹H-NMR (DMSO-d₆) δ ppm: 4.87 (2H, s), 7.57 (1H, dd, J=2.2, 8.8Hz),
5 7.60-7.75 (3H, m), 7.95-8.10 (3H, m), 8.25-8.35 (2H, m), 11.04 (1H, s)

Compound 1-55

¹H-NMR (DMSO-d₆) δ ppm: 2.44 (3H, s), 4.87 (2H, s), 7.30-7.40 (1H, m), 7.45-7.55 (1H, m), 7.60-7.70 (2H, m), 7.90 (1H, brs),
10 7.95-8.05 (3H, m), 8.17 (1H, brs), 10.98 (1H, brs)

Compound 1-56

¹H-NMR (DMSO-d₆) δ ppm: 2.44 (3H, s), 4.89 (2H, s), 7.30-7.40 (1H, m), 7.45-7.80 (3H, m), 7.90 (1H, brs), 7.95-8.15 (3H, m), 8.17 (1H, brs), 10.97 (1H, brs)

15 Compound 1-57

¹H-NMR (DMSO-d₆) δ ppm: 3.87 (3H, s), 4.89 (2H, s), 7.00 (1H, dd, J=2.2, 8.8Hz), 7.11 (1H, d, J=2.2Hz), 7.55-8.15 (6H, m), 8.21 (1H, d, J=8.8Hz), 11.17 (1H, s)

Compound 1-58

20 ¹H-NMR (DMSO-d₆) δ ppm: 3.87 (3H, s), 4.87 (2H, s), 7.00 (1H, dd, J=2.2, 9.1Hz), 7.11 (1H, d, J=2.2Hz), 7.66 (2H, d, J=8.5Hz), 7.75-8.15 (4H, m), 8.24 (1H, d, J=8.8Hz), 11.17 (1H, s)

Compound 1-59

25 ¹H-NMR (DMSO-d₆) δ ppm: 1.43 (3H, t, J=6.9Hz), 4.30 (2H, q, J=6.9Hz), 4.89 (2H, s), 7.10-7.15 (1H, m), 7.20-7.30 (1H, m), 7.55-7.75 (3H, m), 7.91 (1H, brs), 7.95-8.00 (1H, m), 8.05-8.15 (2H, m), 10.95 (1H, s)

Compound 1-60

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.35-7.45 (1H, m), 7.60-7.85 (3H, m), 7.97 (1H, brs), 8.23 (1H, brs), 8.30-8.45 (2H, m), 11.01 (1H, s)

5 Compound 1-61

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.87 (3H, s), 7.00 (1H, dd, $J=2.2, 8.8\text{Hz}$), 7.12 (1H, d, $J=2.2\text{Hz}$), 7.70-7.95 (2H, m), 8.05 (2H, d, $J=8.8\text{Hz}$), 8.30-8.45 (2H, m), 11.15 (1H, s)

Compound 1-62

10 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.87 (3H, s), 7.00 (1H, dd, $J=2.2, 8.8\text{Hz}$), 7.12 (1H, d, $J=2.2\text{Hz}$), 7.75-8.20 (5H, m), 8.29 (1H, d, $J=8.2\text{Hz}$), 8.42 (1H, s), 11.11 (1H, s)

Compound 1-63

15 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.83 (4H, s), 7.35-7.45 (1H, m), 7.55-7.60 (1H, m), 7.65-7.80 (2H, m), 7.90-8.10 (4H, m), 8.27 (1H, brs), 11.02 (1H, s)

Compound 1-64

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.70-2.95 (4H, m), 3.87 (3H, s), 6.90-7.30 (2H, m), 7.40-8.30 (7H, m), 11.16 (1H, s)

20 Compound 1-65

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.23 (3H, s), 2.45 (4H, t, $J=4.7\text{Hz}$), 3.25-3.40 (4H, m), 7.08 (2H, d, $J=8.8\text{Hz}$), 7.20-7.30 (1H, m), 7.52 (1H, dd, $J=2.2, 8.8\text{Hz}$), 7.84 (2H, d, $J=8.8\text{Hz}$), 7.94 (1H, brs), 8.21 (1H, brs), 8.35-8.45 (1H, m), 10.99 (1H, s)

25 Compound 1-66

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.20-3.40 (4H, m), 3.65-3.85 (4H, m), 7.10 (2H, d, $J=9.1\text{Hz}$), 7.20-7.30 (1H, m), 7.53 (1H, dd, $J=2.2,$

8.8Hz), 7.80-8.00 (3H, m), 8.22 (1H, brs), 8.35-8.45 (1H, m),
11.00 (1H, s)

Compound 1-67

¹H-NMR (CDCl₃) δ ppm: 1.40-1.55 (9H, m), 4.35-4.50 (2H, m), 4.98
5 (1H, brs), 5.92 (1H, brs), 6.43 (1H, brs), 7.30-7.50 (4H, m),
7.95-8.05 (2H, m), 8.65-8.75 (1H, m), 10.85 (1H, brs)

Compound 1-68

¹H-NMR (DMSO-d₆) δ ppm: 3.86 (3H, s), 7.20-7.30 (1H, m), 7.35-7.45
(1H, m), 7.48-7.60 (3H, m), 7.67 (1H, dd, J=4.4, 9.1Hz), 7.90-8.10
10 (2H, m), 8.27 (1H, brs), 11.02 (1H, s)

Compound 1-69

¹H-NMR (DMSO-d₆) δ ppm: 3.87 (3H, s), 7.10-7.20 (2H, m), 7.35-7.45
(1H, m), 7.65 (1H, dd, J=4.1, 9.1Hz), 7.90-8.10 (4H, m), 8.26
(1H, brs), 10.95 (1H, s)

15 Compound 1-70

¹H-NMR (DMSO-d₆) δ ppm: 3.86 (3H, s), 7.24 (1H, dd, J=2.5, 7.9Hz),
7.43 (1H, dd, J=1.9, 8.8Hz), 7.48-7.60 (3H, m), 7.76 (1H, d,
J=1.6Hz), 8.02 (1H, brs), 8.20-8.35 (2H, m), 11.06 (1H, s)

Compound 1-71

20 ¹H-NMR (DMSO-d₆) δ ppm: 3.86 (3H, s), 3.87 (3H, s), 6.99 (1H,
dd, J=2.2, 8.8Hz), 7.10 (1H, d, J=2.2Hz), 7.20-7.30 (1H, m),
7.45-7.60 (3H, m), 7.87 (1H, brs), 8.06 (1H, brs), 8.25 (1H,
d, J=9.1Hz), 11.17 (1H, s)

Compound 1-72

25 ¹H-NMR (DMSO-d₆) δ ppm: 3.863 (3H, s), 3.866 (3H, s), 6.98 (1H,
dd, J=2.2, 8.8Hz), 7.09 (1H, d, J=2.2Hz), 7.15-7.20 (2H, m),
7.75-8.15 (4H, m), 8.27 (1H, d, J=8.8Hz), 11.09 (1H, s)

Compound 1-73

¹H-NMR (DMSO-d₆) δ ppm: 1.43 (3H, t, J=6.9Hz), 3.86 (3H, s),
4.30 (2H, q, J=6.9Hz), 7.12 (1H, d, J=7.9Hz), 7.20-7.30 (2H,
m), 7.45-7.65 (3H, m), 7.73 (1H, d, J=8.2Hz), 7.92 (1H, brs),
5 8.09 (1H, brs), 10.95 (1H, s)

Compound 1-74

¹H-NMR (DMSO-d₆) δ ppm: 3.87 (3H, s), 7.10-7.20 (2H, m), 7.35-7.60
(3H, m), 7.69 (1H, dd, J=1.6, 8.5Hz), 7.75-7.85 (3H, m), 7.90-8.05
(3H, m), 8.21 (1H, brs), 8.39 (1H, d, J=8.5Hz), 11.03 (1H, s)
10 Compound 1-75

¹H-NMR (DMSO-d₆) δ ppm: 7.35-7.45 (1H, m), 7.60-7.80 (3H, m),
7.85-8.10 (4H, m), 8.26 (1H, brs), 11.03 (1H, s)

Compound 1-76

¹H-NMR (DMSO-d₆) δ ppm: 7.35-7.45 (1H, m), 7.61 (2H, d, J=7.9Hz),
15 7.67 (1H, dd, J=4.1, 9.1Hz), 7.85-8.05 (2H, m), 8.10-8.35 (3H,
m), 11.00 (1H, s)

Compound 1-77

¹H-NMR (DMSO-d₆) δ ppm: 3.87 (3H, s), 7.00 (1H, dd, J=2.2, 8.8Hz),
7.12 (1H, d, J=2.2Hz), 7.60-8.25 (7H, m), 11.20 (1H, s)
20 Compound 1-78

¹H-NMR (DMSO-d₆) δ ppm: 3.87 (3H, s), 7.00 (1H, dd, J=2.2, 8.8Hz),
7.11 (1H, d, J=2.2Hz), 7.61 (2H, d, J=8.5Hz), 7.70-8.30 (5H,
m), 11.15 (1H, s)

Compound 1-79

¹H-NMR (DMSO-d₆) δ ppm: 1.43 (3H, t, J=7.3Hz), 4.30 (2H, q,
J=7.3Hz), 7.12 (1H, d, J=7.3Hz), 7.20-7.30 (1H, m), 7.62 (1H,
dd, J=0.9, 8.2Hz), 7.68 (1H, d, J=8.2Hz), 7.72-7.80 (1H, m),
25

7.91 (1H, brs), 7.95 (1H, brs), 8.00-8.15 (2H, m), 10.97 (1H, s)

Compound 1-80

¹H-NMR (DMSO-d₆) δ ppm: 7.05-7.50 (7H, m), 7.55-7.80 (4H, m),
5 7.90-7.95 (1H, m), 7.98 (1H, brs), 8.26 (1H, brs), 11.00 (1H, brs)

Compound 1-81

¹H-NMR (DMSO-d₆) δ ppm: 7.05-7.30 (5H, m), 7.35-7.55 (3H, m),
7.60-7.70 (1H, m), 7.90-8.10 (4H, m), 8.26 (1H, brs), 10.96 (1H,
10 s)

Compound 1-82

¹H-NMR (DMSO-d₆) δ ppm: 7.05-7.35 (4H, m), 7.40-7.80 (7H, m),
8.01 (1H, brs), 8.20-8.30 (1H, m), 8.28 (1H, brs), 11.02 (1H, brs)

15 Compound 1-83

¹H-NMR (DMSO-d₆) δ ppm: 7.05-7.35 (4H, m), 7.40-7.65 (5H, m),
7.70-7.80 (2H, m), 8.05 (1H, brs), 8.33 (1H, brs), 10.95 (1H, brs)

Compound 1-84

20 ¹H-NMR (DMSO-d₆) δ ppm: 2.42 (3H, s), 7.05-7.65 (10H, m), 7.75-7.80 (1H, m), 7.88 (1H, brs), 7.90-8.00 (1H, m), 8.16 (1H, brs), 10.97 (1H, brs)

Compound 1-85

¹H-NMR (CDCl₃) δ ppm: 3.88 (3H, s), 5.69 (1H, brs), 6.26 (1H, brs),
25 6.85-7.50 (9H, m), 7.65-7.80 (2H, m), 8.50-8.60 (1H, m), 10.95 (1H, brs)

Compound 1-86

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.10-7.30 (5H, m), 7.35-7.55 (5H, m), 7.65-7.85 (4H, m), 7.97 (1H, brs), 8.00-8.10 (2H, m), 8.22 (1H, brs), 8.35 (1H, d, $J=8.2\text{Hz}$), 11.04 (1H, s)

Compound 1-87

5 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.35-7.60 (4H, m), 7.65-7.85 (4H, m), 7.90-8.10 (4H, m), 8.20-8.35 (2H, m), 11.11 (1H, s)

Compound 1-88

$^1\text{H-NMR}$ ($\text{CDCl}_3+\text{MeOD-d}_4$) δ ppm: 1.60-1.90 (4H, m), 3.00-3.50 (4H, m), 7.25-7.50 (2H, m), 7.80-8.20 (4H, m), 8.45-8.60 (1H, m)

10 Compound 1-89

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 0.75-0.95 (6H, m), 1.40-1.60 (4H, m), 3.00-3.15 (4H, m), 3.87 (3H, s), 7.00 (1H, dd, $J=2.5, 9.1\text{Hz}$), 7.12 (1H, d, $J=2.2\text{Hz}$), 7.87 (1H, brs), 7.95-8.25 (6H, m), 11.21 (1H, s)

15 Compound 1-90

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 6.17 (2H, s), 7.12 (1H, d, $J=8.2\text{Hz}$), 7.35-7.45 (1H, m), 7.50 (1H, d, $J=1.9\text{Hz}$), 7.55-7.70 (2H, m), 7.90-8.05 (2H, m), 8.25 (1H, brs), 10.87 (1H, s)

Compound 1-91

20 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 6.17 (2H, s), 7.12 (1H, d, $J=8.2\text{Hz}$), 7.20-7.30 (1H, m), 7.49 (1H, d, $J=1.6\text{Hz}$), 7.50-7.60 (1H, m), 7.59 (1H, dd, $J=1.6, 8.2\text{Hz}$), 7.94 (1H, brs), 8.20 (1H, brs), 8.25-8.35 (1H, m), 10.94 (1H, s)

Compound 1-92

25 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 6.18 (2H, s), 7.10-7.15 (1H, m), 7.45-7.70 (4H, m), 8.02 (1H, brs), 8.25-8.30 (1H, m), 8.30 (1H, brs), 10.89 (1H, brs)

Compound 1-93

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.87 (3H, s), 6.17 (2H, s), 6.90-7.20 (3H, m), 7.40-7.65 (2H, m), 7.84 (1H, brs), 8.03 (1H, brs), 8.15-8.30 (1H, m), 11.02 (1H, s)

5 Compound 1-94

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.43 (3H, t, $J=6.9\text{Hz}$), 4.29 (2H, q, $J=6.9\text{Hz}$), 6.16 (2H, s), 7.05-7.25 (3H, m), 7.50 (1H, d, $J=1.6\text{Hz}$), 7.59 (1H, dd, $J=1.9, 8.2\text{Hz}$), 7.69 (1H, dd, $J=0.9, 7.9\text{Hz}$), 7.89 (1H, brs), 8.06 (1H, brs), 10.79 (1H, s)

10 Compound 1-95

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.25-3.35 (2H, m), 4.60-4.70 (2H, m), 6.90-7.00 (1H, m), 7.50-7.70 (2H, m), 7.75-7.95 (2H, m), 8.01 (1H, brs), 8.30 (1H, brs), 8.30-8.40 (1H, m), 10.93 (1H, brs)

Compound 1-96

15 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 2.05-2.25 (2H, m), 2.90-3.10 (4H, m), 7.30-7.50 (3H, m), 7.75-7.95 (2H, m), 8.65-8.75 (1H, m)

Compound 1-97

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 3.82 (2H, s), 6.11 (1H, brs), 6.46 (1H, brs), 7.10-7.50 (6H, m), 7.55-7.60 (1H, m), 8.50-8.55 (1H, m),

20 9.94 (1H, brs)

Compound 1-98

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 2.82 (2H, t, $J=7.9\text{Hz}$), 3.11 (2H, t, $J=7.9\text{Hz}$), 5.68 (1H, brs), 6.35 (1H, brs), 7.15-7.55 (7H, m), 8.45-8.55 (1H, m), 9.82 (1H, brs)

25 Compound 1-99

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.81 (3H, s), 3.84 (3H, s), 3.86 (3H, s), 6.90-7.10 (4H, m), 7.20-7.30 (1H, m), 7.35-7.40 (1H, m),

7.55-7.65 (1H, m), 7.76 (1H, brs), 7.94 (1H, brs), 8.05-8.15 (1H, m), 10.26 (1H, brs)

Compound 1-100

¹H-NMR (DMSO-d₆) δ ppm: 3.84 (3H, s), 4.05 (2H, s), 6.93 (1H, dd, J=2.5, 9.1Hz), 6.98-7.15 (3H, m), 7.43 (1H, dd, J=1.3, 5.0Hz), 7.74 (1H, brs), 7.90 (1H, brs), 7.93 (1H, d, J=8.8Hz), 10.34 (1H, s)

Compound 1-101

¹H-NMR (CDCl₃+MeOD-d₄) δ ppm: 6.75-6.90 (1H, m), 7.25-7.75 (4H, m), 7.85-8.00 (1H, m), 8.20-8.30 (1H, m), 8.50-8.65 (1H, m)

Compound 1-102

¹H-NMR (DMSO-d₆) δ ppm: 7.35-7.45 (1H, m), 7.50-7.90 (6H, m), 8.08 (1H, brs), 8.38 (1H, brs), 8.40-8.45 (1H, m), 11.36 (1H, brs)

Compound 1-103

¹H-NMR (CDCl₃+MeOD-d₄) δ ppm: 2.74 (3H, s), 7.25-7.50 (2H, m), 8.30-8.70 (2H, m)

Compound 1-104

¹H-NMR (CDCl₃) δ ppm: 5.64 (1H, brs), 6.41 (1H, brs), 7.10-7.20 (1H, m), 7.35-7.50 (2H, m), 7.60-7.70 (1H, m), 7.75-7.85 (1H, m), 8.65-8.75 (1H, m), 10.84 (1H, brs)

Compound 1-105

¹H-NMR (CDCl₃) δ ppm: 5.64 (1H, brs), 6.40 (1H, brs), 6.80-6.90 (1H, m), 7.30-7.55 (3H, m), 8.10-8.20 (1H, m), 8.65-8.75 (1H, m), 10.54 (1H, brs)

Compound 1-106

¹H-NMR (DMSO-d₆) δ ppm: 1.14 (6H, t, J=7.1Hz), 3.43 (4H, q,

J=7.1Hz), 6.56-6.62 (1H, m), 6.72-6.84 (2H, m), 7.30-7.35 (1H, m), 7.50-8.03 (3H, m), 8.19 (1H, d, J=9.1Hz), 11.21 (1H, s)
Compound 1-107

¹H-NMR (DMSO-d₆) δ ppm: 6.74-6.82 (1H, m), 7.34-7.47 (2H, m),
5 7.62-7.70 (1H, m), 7.96-8.12 (3H, m), 8.31 (1H, brs), 11.04 (1H, s)

Compound 1-108

¹H-NMR (DMSO-d₆) δ ppm: 6.72-6.84 (1H, m), 7.32-7.72 (3H, m),
7.96-8.11 (2H, m), 8.31 (1H, brs), 8.37-8.47 (1H, m), 11.04 (1H,
10 s)

Compound 1-109

¹H-NMR (DMSO-d₆) δ ppm: 6.70-6.90 (1H, m), 7.30-7.70 (2H, m),
7.85-8.20 (3H, m), 8.36 (1H, brs), 10.99 (1H, s)

Compound 1-110

15 ¹H-NMR (DMSO-d₆) δ ppm: 2.43 (3H, s), 6.75-6.80 (1H, m), 7.32-7.39
(2H, m), 7.49 (1H, d, J=8.5Hz), 7.92 (1H, brs), 8.00-8.12 (2H,
m), 8.21 (1H, brs), 10.99 (1H, s)

Compound 1-111

¹H-NMR (DMSO-d₆) δ ppm: 3.81 (3H, s), 6.70-6.85 (1H, m), 7.05-7.60
20 (3H, m), 7.74-8.10 (3H, m), 8.20 (1H, brs), 11.02 (1H, s)

Compound 1-112

¹H-NMR (DMSO-d₆) δ ppm: 3.98 (3H, s), 6.72-6.82 (1H, m), 7.08-7.40
(3H, m), 7.78-8.24 (4H, m), 10.98 (1H, s)

Compound 1-113

25 ¹H-NMR (DMSO-d₆) δ ppm: 3.86 (3H, s), 6.72-7.40 (4H, m), 7.86
(1H, brs), 7.96-8.35 (3H, m), 11.14 (1H, s)

Compound 1-114

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 5.24 (2H, s), 6.72-6.83 (1H, m), 7.00-7.58 (8H, m), 7.85 (1H, brs), 7.97-8.20 (2H, m), 8.30 (1H, d, $J=9.0\text{Hz}$), 11.14 (1H, s)

Compound 1-115

5 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.40 (3H, s), 6.30-6.40 (1H, m), 7.05-7.20 (1H, m), 7.28 (1H, d, $J=3.2\text{Hz}$), 7.45-7.55 (2H, m), 7.86 (1H, brs), 8.06 (1H, brs), 9.98 (1H, s)

Compound 1-116

10 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 2.45 (3H, s), 5.59 (1H, brs), 6.15-6.25 (1H, m), 6.39 (1H, brs), 7.15-7.25 (2H, m), 7.35-7.45 (1H, m), 8.35-8.40 (1H, m), 10.72 (1H, brs)

Compound 1-117

15 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 2.45 (3H, s), 5.59 (1H, brs), 6.15-6.25 (1H, m), 6.38 (1H, brs), 7.15-7.25 (1H, m), 7.35-7.45 (2H, m), 8.65-8.75 (1H, m), 10.73 (1H, brs)

Compound 1-118

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.41 (3H, s), 6.32-6.47 (1H, m), 7.20-7.48 (2H, m), 7.70-7.80 (1H, m), 7.97 (1H, brs), 8.24 (1H, brs), 8.36 (1H, d, $J=8.7\text{Hz}$), 10.99 (1H, s)

20 Compound 1-119

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.41 (3H, s), 6.35-6.47 (1H, m), 7.22-7.33 (1H, m), 7.54-7.64 (1H, m), 7.88-7.96 (1H, m), 8.05 (1H, brs), 8.35 (1H, brs), 10.93 (1H, s)

Compound 1-120

25 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.41 (3H, s), 3.30 (3H, s), 6.35-6.45 (1H, m), 6.95-7.00 (1H, m), 7.05-7.10 (1H, m), 7.20-7.25 (1H, m), 7.82 (1H, brs), 8.03 (1H, brs), 8.25-8.35 (1H, m), 11.07

(1H, brs)

Compound 1-121

¹H-NMR (DMSO-d₆) δ ppm: 1.27 (3H, t, J=7.6Hz), 2.76 (2H, q, J=7.6Hz), 6.40-6.45 (1H, m), 7.25-7.30 (1H, m), 7.35-7.45 (1H, m), 7.60-7.70 (1H, m), 8.00 (1H, brs), 8.10-8.15 (1H, m), 8.27 (1H, brs), 11.06 (1H, brs)

Compound 1-122

¹H-NMR (DMSO-d₆) δ ppm: 1.27 (3H, t, J=7.6Hz), 2.75 (2H, q, J=7.6Hz), 6.35-6.45 (1H, m), 7.20-7.30 (2H, m), 7.53 (1H, dd, J=2.2, 9.1Hz), 7.95 (1H, brs), 8.21 (1H, brs), 8.35-8.50 (1H, m), 11.12 (1H, s)

Compound 1-123

¹H-NMR (DMSO-d₆) δ ppm: 1.27 (3H, t, J=7.6Hz), 2.76 (2H, q, J=7.6Hz), 6.40-6.45 (1H, m), 7.25-7.30 (1H, m), 7.50-7.70 (2H, m), 8.02 (1H, brs), 8.29 (1H, brs), 8.40-8.50 (1H, m), 11.08 (1H, brs)

Compound 1-124

¹H-NMR (DMSO-d₆) δ ppm: 1.26 (3H, t, J=7.5Hz), 2.75 (2H, q, J=7.5Hz), 6.40-6.45 (1H, m), 7.20-7.30 (1H, m), 7.35-7.45 (1H, m), 7.70-7.80 (1H, m), 7.99 (1H, brs), 8.25 (1H, brs), 8.35-8.45 (1H, m), 11.09 (1H, brs)

Compound 1-125

¹H-NMR (DMSO-d₆) δ ppm: 1.26 (3H, t, J=7.8Hz), 2.76 (2H, q, J=7.8Hz), 6.40-6.45 (1H, m), 7.25-7.30 (1H, m), 7.55-7.65 (1H, m), 7.90-8.00 (1H, m), 8.08 (1H, brs), 8.35 (1H, brs), 11.03 (1H, brs)

Compound 1-126

¹H-NMR (DMSO-d₆) δ ppm: 1.27 (3H, t, J=7.6Hz), 2.43 (3H, s), 2.75 (2H, q, J=7.6Hz), 6.40-6.45 (1H, m), 7.20-7.40 (2H, m), 7.45-7.50 (1H, m), 7.90 (1H, brs), 8.10-8.20 (1H, m), 8.17 (1H, brs), 11.01 (1H, brs)

5 Compound 1-127

¹H-NMR (DMSO-d₆) δ ppm: 1.26 (3H, t, J=7.6Hz), 2.75 (2H, q, J=7.6Hz), 3.86 (3H, s), 6.35-6.45 (1H, m), 6.95-7.00 (1H, m), 7.05-7.10 (1H, m), 7.20-7.30 (1H, m), 7.85 (1H, brs), 8.03 (1H, brs), 8.25-8.35 (1H, m), 11.17 (1H, brs)

10 Compound 1-128

¹H-NMR (DMSO-d₆) δ ppm: 1.26 (3H, t, J=7.6Hz), 1.42 (3H, t, J=6.9Hz), 2.75 (2H, q, J=7.6Hz), 4.29 (2H, q, J=6.9Hz), 6.41 (1H, d, J=3.5Hz), 7.12 (1H, d, J=8.2Hz), 7.15-7.30 (2H, m), 7.85 (1H, dd, J=0.9, 8.2Hz), 7.92 (1H, brs), 8.09 (1H, brs), 11.00

15 (1H, s)

Compound 1-129

¹H-NMR (DMSO-d₆) δ ppm: 3.86 (3H, s), 4.93 (2H, s), 6.75-6.85 (1H, m), 6.95-7.15 (2H, m), 7.25-7.35 (1H, m), 7.86 (1H, brs), 8.06 (1H, brs), 8.15-8.30 (1H, m), 11.16 (1H, brs)

20 Compound 1-130

¹H-NMR (DMSO-d₆) δ ppm: 7.25-7.30 (1H, m), 7.40-7.70 (6H, m), 7.90-8.00 (2H, m), 8.15-8.25 (1H, m), 8.16 (1H, brs), 8.36 (1H, brs), 11.48 (1H, brs)

Compound 1-131

25 ¹H-NMR (DMSO-d₆) δ ppm: 7.20-7.30 (1H, m), 7.35-7.55 (5H, m), 7.70-7.80 (1H, m), 7.90-8.00 (2H, m), 8.15 (1H, brs), 8.34 (1H, brs), 8.45-8.55 (1H, m), 11.50 (1H, brs)

Compound 1-132

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.25-7.65 (6H, m), 7.85-8.10 (3H, m),
8.23 (1H, brs), 8.44 (1H, brs), 11.42 (1H, brs)

Compound 1-133

5 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.87 (3H, s), 6.95-7.15 (2H, m), 7.20-7.30
(1H, m), 7.35-7.55 (4H, m), 7.85-8.15 (4H, m), 8.35-8.45 (1H,
m), 11.55 (1H, brs)

Compound 1-134

$^1\text{H-NMR}$ ($\text{CDCl}_3+\text{MeOD-d}_4$) δ ppm: 7.20-7.60 (3H, m), 7.80-8.00 (1H,
10 m), 8.15-8.40 (1H, m), 8.60-8.80 (2H, m)

Compound 1-135

$^1\text{H-NMR}$ ($\text{CDCl}_3+\text{MeOD-d}_4$) δ ppm: 7.30-7.60 (3H, m), 8.20-8.80 (3H,
m), 9.10-9.30 (1H, m)

Compound 1-136

15 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.87 (3H, s), 7.00 (1H, dd, $J=2.2, 8.8\text{Hz}$),
7.12 (1H, d, $J=2.2\text{Hz}$), 7.74-8.20 (4H, m), 8.38 (1H, dd, $J=2.5,$
8.2Hz), 8.99 (1H, d, $J=2.2\text{Hz}$), 11.12 (1H, s)

Compound 1-137

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.15-7.33 (4H, m), 7.35-7.55 (3H, m),
20 7.67 (1H, dd, $J=4.1, 9.1\text{Hz}$), 7.88 (1H, dd, $J=2.8, 9.5\text{Hz}$), 7.98
(1H, brs), 8.24 (1H, brs), 8.40 (1H, dd, $J=2.5, 8.5\text{Hz}$), 8.78
(1H, d, $J=2.5\text{Hz}$), 10.90 (1H, s)

Compound 1-138

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.15-7.35 (4H, m), 7.40-7.60 (4H, m),
25 7.95 (1H, brs), 8.10-8.30 (2H, m), 8.39 (1H, dd, $J=2.5, 8.5\text{Hz}$),
8.77 (1H, d, $J=2.5\text{Hz}$), 10.97 (1H, s)

Compound 1-139

¹H-NMR (DMSO-d₆) δ ppm: 3.45 (3H, s), 4.10 (2H, s), 7.35-7.45 (1H, m), 7.55-7.70 (1H, m), 7.91 (1H, brs), 8.05-8.15 (1H, m), 8.20 (1H, brs), 10.76 (1H, s)

Compound 1-140

5 ¹H-NMR (DMSO-d₆) δ ppm: 3.45 (3H, s), 4.11 (2H, s), 7.54 (1H, dd, J=2.2, 8.8Hz), 7.63 (1H, d, J=8.8Hz), 7.93 (1H, brs), 8.22 (1H, brs), 8.40 (1H, d, J=2.2Hz), 10.76 (1H, s)

Compound 1-141

10 ¹H-NMR (DMSO-d₆) δ ppm: 3.45 (3H, s), 3.96 (3H, s), 4.09 (2H, s), 7.08-7.15 (1H, d, J=7.9Hz), 7.18-7.25 (1H, m), 7.75-7.95 (2H, d, J=8.2Hz), 8.07 (1H, brs), 10.73 (1H, s)

Compound 1-142

15 ¹H-NMR (DMSO-d₆) δ ppm: 4.82 (2H, s), 6.95-7.15 (3H, m), 7.30-7.45 (3H, m), 7.60-7.70 (1H, m), 8.00 (1H, brs), 8.10-8.20 (1H, m), 8.24 (1H, brs), 11.16 (1H, brs)

Compound 1-143

¹H-NMR (CDCl₃) δ ppm: 4.70 (2H, s), 6.90-7.15 (5H, m), 7.30-7.50 (4H, m), 8.55-8.65 (1H, m), 11.17 (1H, brs)

Compound 1-144

20 ¹H-NMR (CDCl₃) δ ppm: 3.88 (3H, s), 4.69 (2H, s), 5.63 (1H, brs), 6.24 (1H, brs), 6.85-7.15 (5H, m), 7.30-7.40 (2H, m), 8.45-8.55 (1H, m), 11.11 (1H, brs)

Compound 1-145

25 ¹H-NMR (DMSO-d₆) δ ppm: 4.83 (2H, s), 7.05-7.15 (2H, m), 7.35-7.45 (3H, m), 7.60-7.70 (1H, m), 8.00 (1H, brs), 8.05-8.15 (1H, m), 8.25 (1H, brs), 11.13 (1H, brs)

Compound 1-146

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 4.84 (2H, s), 7.10-7.15 (2H, m), 7.35-7.45 (2H, m), 7.50-7.70 (2H, m), 8.02 (1H, brs), 8.27 (1H, brs), 8.40-8.45 (1H, m), 11.13 (1H, brs)

Compound 1-147

5 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 3.88 (3H, s), 4.66 (2H, s), 5.54 (1H, brs), 6.26 (1H, brs), 6.80-7.10 (4H, m), 7.20-7.40 (2H, m), 8.45-8.55 (1H, m), 11.09 (1H, brs)

Compound 1-148

10 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.58 (6H, s), 7.05-7.20 (2H, m), 7.30-7.45 (3H, m), 7.60-7.70 (1H, m), 7.95 (1H, brs), 8.05-8.15 (1H, m), 8.22 (1H, brs), 11.24 (1H, brs)

Compound 1-149

15 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.58 (6H, s), 7.05-7.20 (2H, m), 7.30-7.45 (2H, m), 7.50-7.70 (2H, m), 7.97 (1H, brs), 8.24 (1H, brs), 8.40-8.50 (1H, m), 11.28 (1H, brs)

Compound 1-150

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 4.23 (2H, s), 4.70 (2H, s), 7.25-7.45 (4H, m), 7.49 (2H, d, $J=7.3\text{Hz}$), 7.60-7.65 (1H, m), 7.98 (1H, brs), 8.05-8.15 (1H, m), 8.22 (1H, brs), 10.90 (1H, s)

20 Compound 1-151

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 4.23 (2H, s), 4.70 (2H, s), 7.25-7.45 (3H, m), 7.49 (2H, d, $J=7.3\text{Hz}$), 7.55 (1H, dd, $J=2.2, 8.8\text{Hz}$), 7.63 (1H, d, $J=8.8\text{Hz}$), 8.00 (1H, brs), 8.24 (1H, brs), 8.41 (1H, d, $J=2.5\text{Hz}$), 10.90 (1H, s)

25 Compound 1-152

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 0.80-0.95 (4H, m), 1.91-2.05 (1H, m), 7.15-7.25 (1H, m), 7.45-7.53 (1H, m), 7.85 (1H, brs), 7.93-8.02

(1H, m), 8.07 (1H, brs), 10.37 (1H, s)

Compound 1-153

¹H-NMR (DMSO-d₆) δ ppm: 0.82-0.93 (4H, m), 1.94-2.05 (1H, m),
7.31-7.40 (1H, m), 7.67-7.74 (1H, m), 7.82-7.98 (2H, m), 8.11
5 (1H, s), 10.36 (1H, s)

Compound 1-154

¹H-NMR (DMSO-d₆) δ ppm: 0.80-0.95 (4H, m), 1.90-2.00 (1H, m),
7.75-7.95 (1H, m), 7.89 (1H, brs), 8.10 (1H, brs), 10.38 (1H,
brs)

10 Compound 1-155

¹H-NMR (DMSO-d₆) δ ppm: 0.80-1.00 (4H, m), 1.90-2.05 (1H, m),
7.45-7.60 (2H, m), 7.95 (1H, brs), 8.18 (1H, brs), 10.35 (1H,
s)

MS (ESI, m/z): 281 (M+H)⁺

15 Compound 1-156

¹H-NMR (DMSO-d₆) δ ppm: 0.78-0.96 (4H, m), 1.93-2.05 (1H, m),
7.55-7.74 (2H, m), 7.99 (1H, brs), 8.10 (1H, brs), 10.37 (1H,
s)

Compound 1-157

20 ¹H-NMR (DMSO-d₆) δ ppm: 0.81-0.94 (4H, m), 1.90-2.00 (1H, m),
2.44 (3H, s), 7.05-7.15 (1H, m), 7.30-7.40 (1H, m), 7.67-8.10
(3H, m), 10.32 (1H, s)

Compound 1-158

25 ¹H-NMR (DMSO-d₆) δ ppm: 0.80-0.95 (4H, m), 1.90-2.00 (1H, m),
3.31 (3H, s), 7.45-7.55 (1H, m), 7.60-7.70 (1H, m), 7.83 (1H,
brs), 8.06 (1H, brs), 10.34 (1H, brs)

Compound 1-159

¹H-NMR (DMSO-d₆) δ ppm: 0.80-0.90 (4H, m), 1.90-2.00 (1H, m), 3.97 (3H, s), 7.00-7.10 (1H, m), 7.10-7.20 (1H, m), 7.85 (1H, brs), 8.04 (1H, brs), 10.30 (1H, brs)

Compound 1-160

5 ¹H-NMR (DMSO-d₆) δ ppm: 1.75-1.90 (1H, m), 1.90-2.05 (1H, m), 2.10-2.35 (4H, m), 3.20-3.45 (1H, m), 7.30-7.45 (1H, m), 7.55-7.70 (1H, m), 7.75-8.00 (2H, m), 8.15 (1H, brs), 10.05 (1H, s)

MS (ESI, m/z): 277 (M+H)+

10 Compound 1-161

¹H-NMR (DMSO-d₆) δ ppm: 1.75-1.90 (1H, m), 1.90-2.05 (1H, m), 2.10-2.35 (4H, m), 3.30-3.45 (1H, m), 7.50-7.60 (1H, m), 7.60-7.75 (1H, m), 7.95 (1H, brs), 8.19 (1H, brs), 10.03 (1H, s)

15 MS (ESI, m/z): 295 (M+H)+

Compound 1-162

¹H-NMR (DMSO-d₆) δ ppm: 1.50-2.00 (8H, m), 2.85-3.00 (1H, m), 7.30-7.40 (1H, m), 7.55-7.65 (1H, m), 7.75-7.85 (1H, m), 7.87 (1H, brs), 8.11 (1H, brs), 10.14 (1H, brs)

20 Compound 1-163

¹H-NMR (DMSO-d₆) δ ppm: 1.15-1.50 (5H, m), 1.60-2.00 (5H, m), 2.40-2.55 (1H, m), 7.30-7.40 (1H, m), 7.55-7.65 (1H, m), 7.75-7.85 (1H, m), 7.87 (1H, brs), 8.11 (1H, brs), 10.13 (1H, brs)

25 Compound 1-164

¹H-NMR (DMSO-d₆) δ ppm: 1.35-1.45 (1H, m), 1.50-1.60 (1H, m), 2.30-2.55 (2H, m), 7.15-7.40 (5H, m), 7.45-7.60 (2H, m), 7.93

(1H, brs), 8.17 (1H, brs), 10.40 (1H, brs)

Compound 1-165

¹H-NMR (DMSO-d₆) δ ppm: 1.80-2.00 (2H, m), 2.00-2.15 (1H, m),
2.20-2.35 (1H, m), 3.80-3.95 (1H, m), 3.95-4.10 (1H, m),
5 4.45-4.60 (1H, m), 7.30-7.45 (1H, m), 7.55-7.70 (1H, m), 7.91
(1H, brs), 8.05-8.20 (1H, m), 8.21 (1H, brs), 10.90 (1H, s)

Compound 1-166

¹H-NMR (DMSO-d₆) δ ppm: 1.80-2.00 (2H, m), 2.00-2.15 (1H, m),
2.20-2.35 (1H, m), 3.80-3.95 (1H, m), 3.95-4.10 (1H, m),
10 4.45-4.60 (1H, m), 7.30-7.45 (1H, m), 7.55-7.70 (1H, m), 7.91
(1H, brs), 8.05-8.20 (1H, m), 8.21 (1H, brs), 10.90 (1H, s)

Compound 1-167

¹H-NMR (DMSO-d₆) δ ppm: 7.40-7.50 (2H, m), 7.80-7.90 (1H, m),
8.00 (1H, brs), 8.05-8.25 (3H, m), 8.25 (1H, brs), 11.00 (1H,
15 brs)

Compound 1-168

¹H-NMR (DMSO-d₆) δ ppm: 7.40-7.50 (2H, m), 7.55-7.65 (1H, m),
7.70-7.80 (1H, m), 8.05-8.15 (3H, m), 8.37 (1H, brs), 10.92 (1H,
brs)

20 Compound 1-169

¹H-NMR (DMSO-d₆) δ ppm: 2.48 (3H, s), 7.19 (1H, d, J=8.2Hz),
7.42 (1H, s), 7.43-7.60 (1H, m), 7.60-7.73 (1H, m), 7.75-8.00
(3H, m), 8.11 (1H, d, J=8.2Hz), 8.17 (1H, brs), 11.07 (1H, s)

Compound 1-170

25 ¹H-NMR (DMSO-d₆) δ ppm: 2.48 (3H, s), 7.19 (1H, d, J=8.5Hz),
7.35-7.50 (3H, m), 7.89 (1H, brs), 8.00-8.30 (4H, m), 11.04 (1H,
s)

Compound 1-171

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.25 (3H, t, $J=7.6\text{Hz}$), 2.77 (2H, q, $J=7.6\text{Hz}$), 7.22 (1H, d, $J=8.2\text{Hz}$), 7.35-7.50 (3H, m), 7.89 (1H, brs), 8.00-8.25 (4H, m), 11.02 (1H, s)

5 Compound 1-172

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.87 (3H, s), 6.95-7.05 (1H, m), 7.05-7.15 (1H, m), 7.40-7.50 (2H, m), 7.87 (1H, brs), 8.00-8.15 (3H, m), 8.19 (1H, d, $J=8.6\text{Hz}$), 11.12 (1H, s)

Compound 1-173

10 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 6.80-6.95 (2H, m), 7.40-7.50 (2H, m), 7.81 (1H, brs), 8.00-8.10 (3H, m), 8.10-8.15 (1H, m), 10.09 (1H, s), 11.14 (1H, brs)

Compound 1-174

15 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.40-7.50 (2H, m), 7.71 (1H, dd, $J=0.9$, 8.5Hz), 8.00 (1H, s), 8.03-8.20 (3H, m), 8.31 (1H, brs), 8.36 (1H, d, $J=8.5\text{Hz}$), 10.94 (1H, s)

Compound 1-175

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.40 (3H, s), 7.45-7.70 (3H, m), 7.75-8.05 (4H, m), 8.25 (1H, brs), 11.06 (1H, s)

20 Compound 1-176

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.40 (3H, s), 7.35-7.60 (3H, m), 7.85-8.15 (4H, m), 8.25 (1H, brs), 11.04 (1H, s)

Compound 1-177

25 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 4.00 (3H, s), 7.10-7.15 (1H, m), 7.40-7.50 (3H, m), 7.96 (1H, brs), 8.05-8.15 (2H, m), 8.20 (1H, brs), 10.90 (1H, brs)

Compound 1-178

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 6.67-6.84 (1H, m), 7.19-7.53 (3H, m),
7.84-8.15 (4H, m), 10.84 (1H, s), 10.87 (1H, s)

Compound 1-179

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.43 (3H, t, $J=7.0\text{Hz}$), 4.30 (2H, q,
5 $J=7.0\text{Hz}$), 7.07-7.15 (1H, m), 7.19-7.28 (1H, m), 7.56-7.67 (2H,
m), 7.70-7.77 (1H, m), 7.89 (1H, brs), 7.93-8.00 (1H, m),
8.00-8.15 (2H, m), 10.91 (1H, s)

Compound 1-180

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.43 (3H, t, $J=7.0\text{Hz}$), 4.30 (2H, q,
10 $J=7.0\text{Hz}$), 7.08-7.28 (2H, m), 7.60-7.72 (3H, m), 7.90 (1H, brs),
8.00-8.15 (3H, m), 10.93 (1H, s)

Compound 1-181

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 4.00 (3H, s), 7.05-7.15 (1H, m),
7.40-7.45 (1H, m), 7.68 (2H, d, $J=8.5\text{Hz}$), 7.97 (1H, brs), 8.02
15 (2H, d, $J=8.5\text{Hz}$), 8.21 (1H, brs), 10.94 (1H, brs)

Compound 1-182

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.43 (3H, t, $J=7.0\text{Hz}$), 4.30 (2H, q,
 $J=7.0\text{Hz}$), 7.05-7.30 (2H, m), 7.50-7.75 (2H, m), 7.80-7.95 (2H,
m), 8.00-8.15 (2H, m), 10.85 (1H, s)

20 Compound 1-183

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.50-7.75 (3H, m), 8.00-8.10 (2H, m),
8.20-8.30 (1H, m), 8.32 (1H, brs), 10.88 (1H, brs)

Compound 1-184

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.35-7.45 (1H, m), 7.55-7.85 (3H, m),
25 7.97 (1H, brs), 8.00-8.10 (1H, m), 8.23 (1H, brs), 8.30-8.40
(1H, m), 10.90 (1H, brs)

Compound 1-185

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.87 (3H, s), 6.95–7.15 (2H, m), 7.55–7.65 (1H, m), 7.70–7.90 (2H, m), 8.00–8.15 (2H, m), 8.25–8.35 (1H, m), 11.03 (1H, brs)

Compound 1-186

5 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.43 (3H, s), 7.22–7.33 (1H, m), 7.45–7.60 (3H, m), 7.75–7.86 (2H, m), 7.95 (1H, brs), 8.22 (1H, brs), 8.30–8.38 (1H, m), 11.07 (1H, s)

Compound 1-187

10 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.41 (3H, s), 7.22–7.35 (1H, m), 7.35–7.45 (2H, m), 7.50–7.60 (1H, m), 7.85–8.05 (3H, m), 8.22 (1H, brs), 8.32–8.40 (1H, m), 11.08 (1H, s)

Compound 1-188

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.42 (3H, s), 7.40–7.55 (3H, m), 7.75–7.85 (3H, m), 8.02 (1H, brs), 8.25–8.35 (2H, m), 11.04 (1H, brs)

15 Compound 1-189

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.43 (3H, t, $J=7.0\text{Hz}$), 2.41 (3H, s), 4.29 (2H, q, $J=7.0\text{Hz}$), 7.05–7.30 (2H, m), 7.35–7.45 (2H, m), 7.70–7.80 (1H, m), 7.85–8.00 (3H, m), 8.08 (1H, brs), 10.93 (1H, s)

20 Compound 1-190

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.41 (3H, s), 4.00 (3H, s), 7.10–7.15 (1H, m), 7.41 (2H, d, $J=8.0\text{Hz}$), 7.50–7.55 (1H, m), 7.90 (2H, d, $J=8.0\text{Hz}$), 7.97 (1H, brs), 8.22 (1H, brs), 10.95 (1H, brs)

Compound 1-191

25 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.23 (3H, t, $J=7.6\text{Hz}$), 2.71 (2H, q, $J=7.6\text{Hz}$), 4.00 (3H, s), 7.05–7.20 (1H, m), 7.44 (2H, d, $J=8.2\text{Hz}$), 7.50–7.60 (1H, m), 7.92 (2H, d, $J=8.2\text{Hz}$), 7.97 (1H, brs), 8.21

(1H, brs), 10.95 (1H, brs)

Compound 1-192

¹H-NMR (DMSO-d₆) δ ppm: 4.89 (2H, s), 7.20-7.35 (1H, m), 7.50-7.70
 (2H, m), 7.74 (1H, d, J=7.9Hz), 7.85-8.05 (2H, m), 8.09 (1H,
 5 s), 8.15-8.35 (2H, m), 11.09 (1H, s)

Compound 1-193

¹H-NMR (DMSO-d₆) δ ppm: 4.87 (2H, s), 7.20-7.35 (1H, m), 7.50-7.60
 (1H, m), 7.60-7.75 (2H, m), 7.90-8.05 (3H, m), 8.22 (1H, brs),
 8.27-8.35 (1H, m), 11.09 (1H, s)

10 Compound 1-194

¹H-NMR (DMSO-d₆) δ ppm: 4.89 (2H, s), 7.55-7.85 (4H, m), 7.90-8.15
 (3H, m), 8.36 (1H, brs), 10.98 (1H, brs)

Compound 1-195

¹H-NMR (DMSO-d₆) δ ppm: 4.87 (2H, s), 7.50-7.85 (4H, m), 7.95-8.05
 15 (2H, m), 8.09 (1H, brs), 8.36 (1H, brs), 10.98 (1H, brs)

Compound 1-196

¹H-NMR (DMSO-d₆) δ ppm: 2.40 (3H, s), 4.90 (2H, s), 7.50-7.80
 (3H, m), 7.90-8.05 (3H, m), 8.08 (1H, s), 8.26 (1H, brs), 11.10
 (1H, s)

20 Compound 1-197

¹H-NMR (DMSO-d₆) δ ppm: 2.40 (3H, s), 4.88 (2H, s), 7.50-7.75
 (3H, m), 7.90-8.10 (4H, m), 8.26 (1H, brs), 11.10 (1H, s)

Compound 1-198

¹H-NMR (DMSO-d₆) δ ppm: 2.48 (3H, s), 4.89 (2H, s), 7.20 (1H,
 25 d, J=8.5Hz), 7.42 (1H, s), 7.55-7.80 (2H, m), 7.91 (1H, brs),
 7.97 (1H, d, J=7.9Hz), 8.09 (1H, s), 8.15 (1H, d, J=7.9Hz), 8.18
 (1H, brs), 11.10 (1H, s)

Compound 1-199

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.48 (3H, s), 4.87 (2H, s), 7.19 (1H, d, $J=8.2\text{Hz}$), 7.41 (1H, s), 7.66 (2H, d, $J=8.2\text{Hz}$), 7.91 (1H, brs), 8.01 (2H, d, $J=8.2\text{Hz}$), 8.10-8.30 (2H, m), 11.10 (1H, s)

5 Compound 1-200

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 2.50-2.60 (4H, m), 3.61 (2H, s), 3.75-3.85 (4H, m), 3.92 (3H, s), 5.73 (1H, brs), 6.50 (1H, brs), 6.95-7.10 (2H, m), 7.90-8.00 (1H, m), 8.05-8.25 (2H, m), 10.78 (1H, brs)

Compound 1-201

10 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.38 (3H, t, $J=6.9\text{Hz}$), 2.35-2.50 (4H, m), 3.54 (2H, s), 3.55-3.70 (4H, m), 4.16 (2H, q, $J=6.9\text{Hz}$), 7.10-7.25 (1H, m), 7.55-7.65 (1H, m), 7.85-8.05 (3H, m), 8.11 (1H, brs), 8.38 (1H, brs), 10.98 (1H, brs)

Compound 1-202

15 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.35-2.50 (4H, m), 3.55-3.70 (4H, m), 3.58 (2H, s), 5.27 (2H, brs), 7.25-7.55 (7H, m), 7.60-7.70 (1H, m), 7.85-8.10 (4H, m), 8.26 (1H, brs), 11.01 (1H, brs)

Compound 1-203

20 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.35-2.50 (4H, m), 3.57 (2H, s), 3.55-3.70 (4H, m), 5.27 (2H, s), 7.25-7.65 (7H, m), 7.85-8.05 (3H, m), 8.08 (1H, brs), 8.34 (1H, brs), 10.96 (1H, brs)

Compound 1-204

25 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.40-2.55 (4H, m), 2.48 (3H, s), 3.50-3.75 (6H, m), 5.28 (2H, brs), 7.10-7.60 (8H, m), 7.80-8.35 (5H, m), 11.08 (1H, brs)

Compound 1-205

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.40-2.50 (4H, m), 3.55-3.70 (4H, m),

3.58 (2H, s), 3.87 (3H, s), 5.26 (2H, brs), 6.95-7.15 (2H, m),
7.25-7.55 (6H, m), 7.75-8.15 (4H, m), 8.25-8.35 (1H, m), 11.15
(1H, brs)

Compound 1-206

5 $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 7.50-8.50 (8H, m), 10.92 (1H, brs)

Compound 1-207

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 2.48 (3H, s), 3.86 (3H, s), 7.10-7.30
(2H, m), 7.41 (1H, s), 7.45-7.60 (3H, m), 7.90 (1H, brs), 8.10-8.25
(2H, m), 11.10 (1H, s)

10 Compound 1-208

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 2.48 (3H, s), 3.86 (3H, s), 7.14 (2H,
d, $J=8.8\text{Hz}$), 7.18 (1H, d, $J=8.2\text{Hz}$), 7.40 (1H, s), 7.88 (1H, brs),
7.97 (2H, d, $J=8.8\text{Hz}$), 8.15 (1H, brs), 8.21 (1H, d, $J=8.2\text{Hz}$),
11.02 (1H, s)

15 Compound 1-209

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 3.86 (3H, s), 7.20-7.35 (2H, m), 7.45-7.65
(4H, m), 7.96 (1H, brs), 8.22 (1H, brs), 8.29-8.39 (1H, m), 11.09
(1H, s)

Compound 1-210

20 $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 3.86 (3H, s), 7.20-7.30 (1H, m), 7.45-7.65
(4H, m), 7.75-7.90 (1H, m), 8.09 (1H, brs), 8.37 (1H, brs), 10.97
(1H, brs)

Compound 1-211

25 $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 3.87 (3H, s), 7.10-7.20 (2H, m), 7.50-7.65
(1H, m), 7.80-8.05 (3H, m), 8.06 (1H, brs), 8.33 (1H, brs), 10.90
(1H, brs)

Compound 1-212

¹H-NMR (DMSO-d₆) δ ppm: 2.40 (3H, s), 3.86 (3H, s), 7.20-7.30 (1H, m), 7.45-7.65 (4H, m), 7.90-8.10 (2H, m), 8.26 (1H, brs), 11.10 (1H, s)

Compound 1-213

5 ¹H-NMR (DMSO-d₆) δ ppm: 3.86 (3H, s), 3.99 (3H, s), 7.05-7.15 (3H, m), 7.50-7.55 (1H, m), 7.90-8.00 (3H, m), 8.20 (1H, brs), 10.89 (1H, brs)

Compound 1-214

10 ¹H-NMR (DMSO-d₆) δ ppm: 1.32 (6H, d, J=6.0Hz), 4.65-4.80 (1H, m), 7.15-7.25 (1H, m), 7.35-7.60 (4H, m), 7.63-7.70 (1H, m), 7.90-8.10 (2H, m), 8.27 (1H, brs), 11.01 (1H, s)

Compound 1-215

15 ¹H-NMR (DMSO-d₆) δ ppm: 1.01 (3H, t, J=7.6Hz), 1.70-1.85 (2H, dt, J=6.6, 7.6Hz), 4.03 (2H, t, J=6.6Hz), 7.20-7.25 (1H, m), 7.35-7.60 (4H, m), 7.63-7.70 (1H, m), 7.90-8.10 (2H, m), 8.27 (1H, brs), 11.01 (1H, s)

Compound 1-216

20 ¹H-NMR (DMSO-d₆) δ ppm: 7.23-7.33 (1H, m), 7.50-7.60 (1H, m), 7.65-7.80 (2H, m), 7.90-8.08 (3H, m), 8.15-8.30 (2H, m), 11.11 (1H, s)

Compound 1-217

¹H-NMR (DMSO-d₆) δ ppm: 1.43 (3H, t, J=7.0Hz), 4.30 (2H, q, J=7.0Hz), 7.09-7.28 (2H, m), 7.55-7.68 (3H, m), 7.90 (1H, brs), 8.00-8.20 (3H, m), 10.94 (1H, s)

25 Compound 1-218

¹H-NMR (DMSO-d₆) δ ppm: 4.00 (3H, s), 7.05-7.15 (1H, m), 7.35-7.45 (1H, m), 7.60 (2H, d, J=8.5Hz), 7.96 (1H, brs), 8.14 (2H, d,

J=8.5Hz), 8.20 (1H, brs), 10.94 (1H, brs)

Compound 1-219

¹H-NMR (DMSO-d₆) δ ppm: 3.33 (3H, s), 3.65-3.75 (2H, m), 4.15-4.25
(2H, m), 7.20-7.30 (1H, m), 7.35-7.70 (5H, m), 7.90-8.10 (2H,
5 m), 8.27 (1H, brs), 11.00 (1H, s)

Compound 1-220

¹H-NMR (DMSO-d₆) δ ppm: 3.31 (3H, s), 3.33 (3H, s), 3.65-3.75
(2H, m), 4.10-4.30 (2H, m), 7.25 (1H, dd, J=2.2, 7.6Hz), 7.35-7.75
(4H, m), 7.90-8.05 (2H, m), 8.26 (1H, brs), 11.00 (1H, s)

10 Compound 1-221

¹H-NMR (CDCl₃) δ ppm: 1.20-1.65 (6H, m), 1.75-1.90 (2H, m),
1.95-2.10 (2H, m), 4.30-4.45 (1H, m), 5.69 (1H, brs), 6.41 (1H,
brs), 7.05-7.65 (6H, m), 8.35-8.45 (1H, m), 10.82 (1H, brs)

Compound 1-222

15 ¹H-NMR (CDCl₃) δ ppm: 1.20-1.65 (6H, m), 1.75-1.90 (2H, m),
1.95-2.10 (2H, m), 2.51 (3H, s), 4.30-4.45 (1H, m), 5.55 (1H,
brs), 6.36 (1H, brs), 7.05-7.30 (3H, m), 7.35-7.45 (1H, m),
7.50-7.65 (2H, m), 8.50-8.60 (1H, m), 10.90 (1H, brs)

Compound 1-223

20 ¹H-NMR (CDCl₃) δ ppm: 1.75-1.95 (2H, m), 2.00-2.15 (2H, m),
3.55-3.70 (2H, m), 3.95-4.05 (2H, m), 4.55-4.70 (1H, m), 6.08
(1H, brs), 6.48 (1H, brs), 7.10-7.65 (6H, m), 8.30-8.45 (1H,
m), 10.86 (1H, brs)

Compound 1-224

25 ¹H-NMR (CDCl₃) δ ppm: 1.75-1.90 (2H, m), 2.00-2.15 (2H, m), 2.51
(3H, s), 3.55-3.65 (2H, m), 3.95-4.05 (2H, m), 4.55-4.70 (1H,
m), 5.60 (1H, brs), 6.39 (1H, brs), 7.10-7.30 (3H, m), 7.35-7.50

(1H, m), 7.55-7.65 (2H, m), 8.45-8.55 (1H, m), 10.94 (1H, brs)

Compound 1-225

¹H-NMR (CDCl₃) δ ppm: 1.80-2.15 (4H, m), 2.25-2.45 (2H, m), 2.33
(3H, s), 2.65-2.80 (2H, m), 4.40-4.55 (1H, m), 6.00 (1H, brs),
5 6.48 (1H, brs), 7.05-7.30 (2H, m), 7.35-7.65 (4H, m), 8.30-8.45
(1H, m), 10.85 (1H, brs)

Compound 1-226

¹H-NMR (DMSO-d₆) δ ppm: 1.60-1.75 (2H, m), 1.90-2.05 (2H, m),
2.10-2.30 (5H, m), 2.55-2.75 (2H, m), 4.40-4.55 (1H, m),
10 7.20-7.30 (1H, m), 7.40-7.60 (4H, m), 7.70-7.80 (1H, m), 8.00
(1H, brs), 8.15-8.35 (2H, m), 11.02 (1H, s)

MS (ESI, m/z): 428 (M+H)+

Compound 1-227

¹H-NMR (CDCl₃) δ ppm: 1.80-2.15 (4H, m), 2.20-2.45 (2H, m), 2.33
15 (3H, s), 2.60-2.80 (2H, m), 4.40-4.50 (1H, m), 5.96 (1H, brs),
6.56 (1H, brs), 7.00-7.20 (2H, m), 7.35-7.65 (3H, m), 8.15-8.25
(1H, m), 10.81 (1H, brs)

Compound 1-228

¹H-NMR (CDCl₃) δ ppm: 1.80-2.10 (4H, m), 2.25-2.40 (2H, m), 2.32
20 (3H, s), 2.50 (3H, s), 2.65-2.80 (2H, m), 4.40-4.50 (1H, m),
5.81 (1H, brs), 6.41 (1H, brs), 7.00-7.30 (3H, m), 7.35-7.65
(3H, m), 8.45-8.55 (1H, m), 10.92 (1H, brs)

Compound 1-229

¹H-NMR (DMSO-d₆) δ ppm: 3.23-3.36 (2H, m), 4.60-4.70 (2H, m),
25 6.95 (1H, d, J=8.2Hz), 7.53-7.63 (1H, m), 7.77-7.92 (3H, m),
8.05 (1H, brs), 8.33 (1H, brs), 10.85 (1H, s)

Compound 1-230

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.47 (3H, s), 6.17 (2H, s), 7.12 (1H, d, $J=8.2\text{Hz}$), 7.18 (1H, d, $J=8.2\text{Hz}$), 7.40 (1H, s), 7.48 (1H, d, $J=1.6\text{Hz}$), 7.58 (1H, dd, $J=1.6, 8.2\text{Hz}$), 7.88 (1H, brs), 8.05-8.25 (2H, m), 10.95 (1H, s)

5 Compound 1-231

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.25 (3H, t, $J=7.6\text{Hz}$), 2.77 (2H, q, $J=7.6\text{Hz}$), 6.17 (2H, s), 7.05-7.15 (1H, m), 7.22 (1H, d, $J=8.2\text{Hz}$), 7.41 (1H, s), 7.49 (1H, s), 7.58 (1H, d, $J=8.2\text{Hz}$), 7.88 (1H, brs), 8.13 (1H, brs), 8.17 (1H, d, $J=8.2\text{Hz}$), 10.93 (1H, s)

10 Compound 1-232

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.39 (3H, s), 6.17 (2H, s), 7.05-7.20 (1H, m), 7.48 (1H, d, $J=1.9\text{Hz}$), 7.52-7.60 (2H, m), 7.90-8.05 (2H, m), 8.24 (1H, brs), 10.95 (1H, s)

Compound 1-233

15 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.68 (3H, s), 7.35-7.45 (1H, m), 7.67 (1H, dd, $J=4.1, 9.1\text{Hz}$), 7.87 (1H, d, $J=8.5\text{Hz}$), 7.93-8.10 (3H, m), 8.26 (1H, brs), 8.30 (1H, d, $J=1.6\text{Hz}$), 11.03 (1H, s)

Compound 1-234

20 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.92 (3H, s), 7.35-7.48 (1H, m), 7.64-7.95 (3H, m), 8.02 (1H, brs), 8.20-8.35 (3H, m), 8.55-8.65 (1H, m), 11.11 (1H, s)

Compound 1-235

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 5.43 (2H, s), 7.32-8.08 (10H, m), 8.22-8.35 (3H, m), 8.59-8.65 (1H, m), 11.12 (1H, s)

25 Compound 1-236

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.68 (3H, s), 7.35-7.45 (1H, m), 7.65-7.80 (1H, m), 7.90-8.00 (1H, m), 8.01 (1H, brs), 8.20-8.30 (3H, m),

8.27 (1H, brs), 8.55-8.60 (1H, m), 11.12 (1H, brs)

Compound 1-237

¹H-NMR (DMSO-d₆) δ ppm: 2.48 (3H, s), 2.68 (3H, s), 7.15-7.25
(1H, m), 7.40-7.45 (1H, m), 7.70-7.80 (1H, m), 7.91 (1H, brs),
5 8.10-8.15 (1H, m), 8.17 (1H, brs), 8.20-8.30 (2H, m), 8.50-8.60
(1H, m), 11.19 (1H, brs)

Compound 1-238

¹H-NMR (DMSO-d₆) δ ppm: 1.60-1.75 (2H, m), 1.90-2.05 (2H, m),
2.20-2.35 (2H, m), 2.60-2.75 (2H, m), 3.50 (2H, s), 4.45-4.60
10 (1H, m), 7.20-7.70 (11H, m), 7.95-8.05 (1H, m), 8.02 (1H, brs),
8.30 (1H, brs), 10.98 (1H, brs)

Compound 1-239

¹H-NMR (DMSO-d₆) δ ppm: 1.60-1.75 (2H, m), 1.90-2.05 (2H, m),
2.20-2.35 (2H, m), 2.60-2.75 (2H, m), 3.50 (2H, s), 4.45-4.60
15 (1H, m), 7.20-7.60 (10H, m), 7.70-7.80 (1H, m), 7.99 (1H, brs),
8.15-8.35 (2H, m), 11.01 (1H, s)

MS (ESI, m/z): 504 (M+H)⁺

Compound 1-240

¹H-NMR (DMSO-d₆) δ ppm: 1.60-1.80 (2H, m), 1.90-2.05 (2H, m),
20 2.20-2.35 (2H, m), 2.60-2.80 (2H, m), 3.50 (2H, s), 4.45-4.60
(1H, m), 7.20-7.35 (6H, m), 7.40-7.65 (4H, m), 7.75-7.85 (1H,
m), 8.07 (1H, brs), 8.33 (1H, brs), 10.92 (1H, brs)

Compound 1-241

¹H-NMR (DMSO-d₆) δ ppm: 1.60-1.75 (2H, m), 1.90-2.05 (2H, m),
25 2.20-2.35 (2H, m), 2.48 (3H, s), 2.60-2.75 (2H, m), 3.50 (2H,
s), 4.45-4.60 (1H, m), 7.15-7.60 (10H, m), 7.91 (1H, brs),
8.15-8.20 (1H, m), 8.19 (1H, brs), 11.06 (1H, brs)

Compound 1-242

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.43 (3H, t, $J=7.0\text{Hz}$), 4.29 (2H, q, $J=7.0\text{Hz}$), 7.05-7.50 (8H, m), 7.55-7.82 (4H, m), 7.90 (1H, brs), 8.08 (1H, brs), 10.93 (1H, s)

5 Compound 1-243

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.43 (3H, t, $J=7.0\text{Hz}$), 4.29 (2H, q, $J=7.0\text{Hz}$), 7.05-7.30 (7H, m), 7.40-7.52 (2H, m), 7.65-7.75 (1H, m), 7.90 (1H, brs), 8.00-8.15 (3H, m), 10.88 (1H, s)

Compound 1-244

10 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.47 (3H, s), 7.05-7.25 (4H, m), 7.31 (1H, dd, $J=2.2, 8.2\text{Hz}$), 7.35-7.50 (3H, m), 7.53-7.65 (2H, m), 7.70-7.80 (1H, m), 7.88 (1H, brs), 8.12 (1H, d, $J=8.2\text{Hz}$), 8.16 (1H, brs), 11.08 (1H, s)

Compound 1-245

15 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.48 (3H, s), 7.10-7.30 (6H, m), 7.35-7.55 (3H, m), 7.89 (1H, brs), 7.95-8.07 (2H, m), 8.10-8.25 (2H, m), 11.03 (1H, s)

Compound 1-246

20 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.48 (3H, s), 7.00-7.10 (2H, m), 7.15-7.25 (1H, m), 7.35-7.55 (2H, m), 7.65-7.80 (2H, m), 7.90 (1H, brs), 7.90-8.00 (1H, m), 8.05-8.15 (1H, m), 8.18 (1H, brs), 8.45-8.55 (2H, m), 11.08 (1H, brs)

Compound 1-247

25 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.83 (4H, s), 7.50-7.80 (4H, m), 7.90-8.10 (3H, m), 8.20-8.35 (2H, m), 11.03 (1H, s)

Compound 1-248

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.82 (2H, s), 7.25-7.45 (6H, m), 7.55-7.75

(2H, m), 7.86 (1H, brs), 8.08 (1H, brs), 10.25 (1H, brs)

Compound 1-249

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 3.82 (2H, s), 7.20-7.60 (7H, m), 7.94 (1H, brs), 8.16 (1H, brs), 10.27 (1H, s)

5 Compound 1-250

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 1.40-1.55 (3H, m), 3.95-4.10 (1H, m), 7.15-7.60 (7H, m), 7.93 (1H, brs), 8.15 (1H, brs), 10.26 (1H, brs)

Compound 1-251

10 $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 1.85-2.00 (2H, m), 2.40-2.73 (4H, m), 7.15-7.40 (6H, m), 7.56-7.75 (2H, m), 7.88 (1H, brs), 8.11 (1H, brs), 10.10 (1H, s)

Compound 1-252

15 $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 1.87-2.02 (2H, m), 2.40-2.73 (7H, m), 7.08-7.44 (7H, m), 7.78 (1H, brs), 7.88-7.96 (1H, m), 8.00 (1H, brs), 10.11 (1H, s)

Compound 1-253

20 $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 1.60-1.70 (4H, m), 2.40-2.67 (4H, m), 7.11-7.41 (6H, m), 7.56-7.75 (2H, m), 7.88 (1H, brs), 8.11 (1H, brs), 10.10 (1H, s)

Compound 1-254

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 1.60-1.70 (4H, m), 2.40-2.67 (7H, m), 7.08-7.40 (7H, m), 7.77 (1H, brs), 7.86-7.94 (1H, m), 8.00 (1H, brs), 10.10 (1H, s)

25 Compound 1-255

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 3.77 (3H, s), 3.78 (2H, s), 6.80-7.05 (3H, m), 7.20-7.35 (2H, m), 7.55-7.75 (2H, m), 7.87 (1H, brs),

8.09 (1H, brs), 10.24 (1H, brs)

Compound 1-256

¹H-NMR (DMSO-d₆) δ ppm: 3.73 (2H, s), 3.74 (3H, s), 6.85-6.95 (2H, m), 7.25-7.40 (3H, m), 7.55-7.75 (2H, m), 7.86 (1H, brs),

5 8.08 (1H, brs), 10.20 (1H, brs)

Compound 1-257

¹H-NMR (DMSO-d₆) δ ppm: 1.84-1.95 (2H, m), 2.35-2.55 (5H, m), 2.59 (2H, t, J=7.7Hz), 3.71 (3H, s), 6.80-6.90 (2H, m), 7.10-7.20 (3H, m), 7.36 (1H, s), 7.77 (1H, brs), 7.90 (1H, d, J=8.2Hz),

10 8.00 (1H, brs), 10.09 (1H, s)

Compound 1-258

¹H-NMR (DMSO-d₆) δ ppm: 1.86-1.98 (2H, m), 2.40-2.47 (5H, m), 2.59 (2H, t, J=7.5Hz), 3.71 (3H, s), 3.74 (3H, s), 6.70-6.90 (3H, m), 7.10-7.20 (1H, m), 7.36 (1H, s), 7.77 (1H, brs), 7.92

15 (1H, d, J=8.1Hz), 8.01 (1H, brs), 10.10 (1H, s)

Compound 1-259

¹H-NMR (DMSO-d₆) δ ppm: 4.13 (2H, s), 6.89 (1H, s), 7.10-7.80 (7H, m), 7.92 (1H, brs), 8.14 (1H, brs), 10.44 (1H, brs)

Compound 1-260

20 ¹H-NMR (DMSO-d₆) δ ppm: 2.45 (3H, s), 4.12 (2H, s), 6.88 (1H, s), 7.10-7.65 (6H, m), 7.80 (1H, brs), 7.88-7.96 (1H, m), 8.04 (1H, brs), 10.43 (1H, brs)

Compound 1-261

¹H-NMR (DMSO-d₆) δ ppm: 1.92-2.03 (2H, m), 2.47-2.55 (2H, m), 2.89 (2H, t, J=7.5Hz), 6.85-7.00 (2H, m), 7.30-7.40 (2H, m), 7.58-7.75 (2H, m), 7.89 (1H, brs), 8.11 (1H, brs), 10.11 (1H,

s)

Compound 1-262

¹H-NMR (DMSO-d₆) δ ppm: 1.90-2.04 (2H, m), 2.45 (3H, s), 2.47-2.53 (2H, m), 2.89 (2H, t, J=7.6Hz), 6.86-7.00 (2H, m), 7.13 (1H, d, J=8.1Hz), 7.30-7.40 (2H, m), 7.77 (1H, brs), 7.91 (1H, d, J=8.3Hz), 8.00 (1H, brs), 10.11 (1H, s)

Compound 1-263

¹H-NMR (DMSO-d₆) δ ppm: 1.50 (3H, d, J=6.9Hz), 2.18 (3H, s), 5.22 (1H, q, J=6.9Hz), 7.50-7.65 (1H, m), 7.65-7.80 (1H, m), 8.07 (1H, brs), 8.29 (1H, brs), 10.71 (1H, s)

10 Compound 1-264

¹H-NMR (DMSO-d₆) δ ppm: 3.45 (3H, s), 4.11 (2H, s), 7.50-7.65 (1H, m), 7.90-8.10 (2H, m), 8.29 (1H, brs), 10.74 (1H, s)

Compound 1-265

¹H-NMR (DMSO-d₆) δ ppm: 1.60 (3H, d, J=6.7Hz), 5.10 (1H, q, J=6.7Hz), 6.95-7.15 (3H, m), 7.25-7.40 (2H, m), 7.50-7.65 (1H, m), 7.80-7.95 (1H, m), 8.05 (1H, brs), 8.30 (1H, brs), 11.10 (1H, brs)

Compound 1-266

¹H-NMR (DMSO-d₆) δ ppm: 2.85-3.00 (2H, m), 4.25-4.40 (2H, m), 6.90-7.05 (2H, m), 7.20-7.70 (6H, m), 7.89 (1H, brs), 8.08 (1H, brs), 10.27 (1H, brs)

Compound 1-267

¹H-NMR (DMSO-d₆) δ ppm: 2.85-3.00 (2H, m), 4.20-4.40 (2H, m), 6.90-7.05 (3H, m), 7.20-7.45 (3H, m), 7.65-7.80 (1H, m), 7.90-8.05 (1H, m), 7.91 (1H, brs), 8.09 (1H, brs), 10.31 (1H, brs)

Compound 1-268

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.02-2.14 (2H, m), 2.45 (3H, s), 2.58-2.65 (2H, m), 3.98-4.10 (2H, m), 6.86-6.98 (3H, m), 7.08-7.38 (4H, m), 7.78 (1H, brs), 7.91 (1H, d, $J=8.1\text{Hz}$), 7.91 (1H, brs), 10.15 (1H, s)

5 Compound 1-269

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.03-2.13 (2H, m), 2.63 (2H, t, $J=7.4\text{Hz}$), 4.05 (2H, t, $J=6.4\text{Hz}$), 6.87-7.00 (3H, m), 7.22-7.40 (3H, m), 7.57-7.77 (2H, m), 7.89 (1H, brs), 8.11 (1H, brs), 10.15 (1H, s)

10 Compound 1-270

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.92 (2H, t, $J=6.0\text{Hz}$), 4.28 (2H, t, $J=6.0\text{Hz}$), 6.95-7.05 (2H, m), 7.08-7.18 (2H, m), 7.30-7.40 (1H, m), 7.55-7.70 (2H, m), 7.91 (1H, brs), 8.11 (1H, brs), 10.28 (1H, s)

15 Compound 1-271

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.85 (3H, s), 4.22 (2H, s), 4.69 (2H, s), 6.90-7.00 (1H, m), 7.02-7.10 (1H, m), 7.28-7.42 (3H, m), 7.46-7.54 (2H, m), 7.82 (1H, brs), 7.97 (1H, brs), 8.30 (1H, d, $J=9.0\text{Hz}$), 10.98 (1H, s)

20 Compound 1-272

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.83 (3H, s), 4.05 (2H, s), 6.85-6.95 (1H, m), 7.00-7.10 (1H, m), 7.15-7.50 (5H, m), 7.77 (1H, brs), 7.85-8.00 (2H, m), 10.84 (1H, brs)

Compound 1-273

25 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 4.06 (2H, s), 7.15-7.60 (7H, m), 8.00 (1H, brs), 8.23 (1H, brs), 10.74 (1H, brs)

Compound 1-274

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.19–3.88 (8H, m), 7.36–7.48 (1H, m), 7.63–7.77 (3H, m), 7.88–8.16 (4H, m), 8.30 (1H, s), 11.03 (1H, s)

Compound 1-275

5 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.10–2.60 (7H, m), 3.10–3.80 (4H, m), 7.52–7.80 (4H, m), 7.95–8.15 (3H, m), 8.32 (1H, brs), 10.97 (1H, s)

MS (ESI, m/z): 443 ($\text{M}+\text{H}$) $^+$

Compound 1-276

10 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.15 (3H, s), 2.30–2.45 (4H, m), 2.90–3.05 (4H, m), 7.55–7.65 (1H, m), 7.70–7.80 (1H, m), 7.85–8.05 (2H, m), 8.06 (1H, brs), 8.25–8.40 (3H, m), 11.09 (1H, brs)

Compound 1-277

15 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.40–2.50 (4H, m), 2.90–3.05 (4H, m), 3.47 (2H, s), 7.15–7.35 (5H, m), 7.50–7.65 (1H, m), 7.70–7.80 (1H, m), 7.85–8.05 (2H, m), 8.06 (1H, brs), 8.25–8.40 (3H, m), 11.09 (1H, brs)

Compound 1-278

20 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.48 (3H, s), 7.15–7.45 (3H, m), 7.95 (1H, brs), 8.05–8.15 (1H, m), 8.22 (1H, brs), 8.80–8.90 (1H, m), 11.34 (1H, brs)

Compound 1-279

25 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.25–7.50 (2H, m), 7.60–7.75 (1H, m), 7.90–8.00 (1H, m), 8.02 (1H, brs), 8.35 (1H, brs), 8.80–8.95 (1H, m), 11.27 (1H, brs)

Compound 1-280

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.25–7.45 (2H, m), 7.60–7.70 (1H, m),

7.85-8.00 (3H, m), 8.00 (1H, brs), 8.26 (1H, brs), 10.94 (1H, brs)

Compound 1-281

¹H-NMR (DMSO-d₆) δ ppm: 6.85-6.95 (1H, m), 7.30-7.50 (2H, m),
5 7.60-7.70 (1H, m), 7.95-8.05 (2H, m), 8.28 (1H, brs), 11.00 (1H, brs)

Compound 1-282

¹H-NMR (DMSO-d₆) δ ppm: 1.26 (3H, t, J=7.6Hz), 2.47 (3H, s),
2.75 (2H, q, J=7.6Hz), 6.35-6.45 (1H, m), 7.18 (1H, d, J=8.2Hz),
10 7.24 (1H, d, J=3.5Hz), 7.39 (1H, s), 7.89 (1H, brs), 8.16 (1H, brs), 8.27 (1H, d, J=8.2Hz), 11.11 (1H, s)

Compound 1-283

¹H-NMR (DMSO-d₆) δ ppm: 1.20-1.30 (6H, m), 2.70-2.82 (4H, m),
6.41 (1H, d, J=3.2Hz), 7.21 (1H, dd, J=1.6, 8.2Hz), 7.25 (1H,
15 d, J=3.2Hz), 7.40 (1H, s), 7.89 (1H, brs), 8.14 (1H, brs), 8.29 (1H, d, J=8.2Hz), 11.10 (1H, s)

Compound 1-284

¹H-NMR (DMSO-d₆) δ ppm: 1.20-1.30 (9H, m), 2.75 (2H, q, J=7.6Hz),
3.00-3.15 (1H, m), 6.35-6.45 (1H, m), 7.20-7.30 (2H, m),
20 7.35-7.45 (1H, m), 7.90 (1H, brs), 8.12 (1H, brs), 8.29 (1H, d, J=8.5Hz), 11.08 (1H, s)

Compound 1-285

¹H-NMR (DMSO-d₆) δ ppm: 1.27 (3H, t, J=7.6Hz), 1.36 (9H, s),
2.76 (2H, q, J=7.6Hz), 6.42 (1H, d, J=3.2Hz), 7.26 (1H, d,
25 J=3.2Hz), 7.44 (1H, dd, J=1.9, 8.5Hz), 7.48 (1H, d, J=1.9Hz),
7.93 (1H, brs), 8.13 (1H, brs), 8.30 (1H, d, J=8.5Hz), 11.10 (1H, s)

Compound 1-286

^1H -NMR (DMSO- d_6) δ ppm: 1.27 (3H, t, $J=7.6\text{Hz}$), 2.76 (2H, q, $J=7.6\text{Hz}$), 6.40-6.45 (1H, m), 7.29 (1H, d, $J=3.5\text{Hz}$), 7.70 (1H, dd, $J=1.3, 8.5\text{Hz}$), 7.97 (1H, s), 8.10 (1H, brs), 8.33 (1H, brs),
5 8.55 (1H, d, $J=8.5\text{Hz}$), 11.02 (1H, s)

Compound 1-287

^1H -NMR (DMSO- d_6) δ ppm: 1.26 (3H, t, $J=7.6\text{Hz}$), 1.37 (3H, t, $J=6.9\text{Hz}$), 2.75 (2H, q, $J=7.6\text{Hz}$), 4.13 (2H, q, $J=6.9\text{Hz}$), 6.41 (1H, d, $J=3.2\text{Hz}$), 6.96 (1H, dd, $J=2.2, 8.8\text{Hz}$), 7.06 (1H, d, $J=2.2\text{Hz}$),
10 7.24 (1H, d, $J=3.2\text{Hz}$), 7.84 (1H, brs), 8.03 (1H, brs), 8.31 (1H, d, $J=8.8\text{Hz}$), 11.18 (1H, s)

Compound 1-288

^1H -NMR (CDCl_3) δ ppm: 1.34 (3H, t, $J=7.6\text{Hz}$), 2.79 (2H, q, $J=7.6\text{Hz}$), 5.67 (1H, brs), 6.15-6.25 (1H, m), 6.34 (1H, brs), 7.20-7.30 (2H, m),
15 8.55-8.65 (1H, m), 10.80 (1H, brs)

Compound 1-289

^1H -NMR (DMSO- d_6) δ ppm: 1.26 (3H, t, $J=7.6\text{Hz}$), 2.30-2.45 (3H, m), 2.75 (2H, q, $J=7.6\text{Hz}$), 6.42 (1H, d, $J=3.5\text{Hz}$), 7.26 (1H, d, $J=3.5\text{Hz}$), 7.54 (1H, d, $J=3.5\text{Hz}$), 7.97 (1H, brs), 8.10 (1H, d, $J=10.4\text{Hz}$),
20 8.25 (1H, brs), 11.13 (1H, s)

Compound 1-290

^1H -NMR (DMSO- d_6) δ ppm: 1.26 (3H, t, $J=7.5\text{Hz}$), 2.75 (2H, q, $J=7.5\text{Hz}$), 3.99 (3H, s), 6.35-6.45 (1H, m), 7.05-7.30 (1H, m), 7.60-7.70 (1H, m), 7.99 (1H, brs), 8.23 (1H, brs), 11.03 (1H, brs)
25

Compound 1-291

^1H -NMR (DMSO- d_6) δ ppm: 1.34 (9H, s), 6.41 (1H, d, $J=3.5\text{Hz}$),

7.26 (1H, d, J=3.5Hz), 7.55-7.70 (1H, m), 8.04-8.10 (1H, m),
8.19 (1H, brs), 8.41 (1H, brs), 11.30 (1H, s)

Compound 1-292

¹H-NMR (DMSO-d₆) δ ppm: 0.82-0.95 (3H, m), 1.27-1.39 (4H, m),
5 1.62-1.75 (2H, m), 2.73 (2H, t, J=7.6Hz), 6.43 (1H, d, J=3.2Hz),
7.22-7.46 (2H, m), 7.60-7.70 (1H, m), 8.00 (1H, brs), 8.08-8.16
(1H, m), 8.27 (1H, brs), 11.03 (1H, s)

Compound 1-293

¹H-NMR (DMSO-d₆) δ ppm: 0.82-0.95 (3H, m), 1.25-1.40 (4H, m),
10 1.60-1.80 (2H, m), 2.47 (3H, s), 2.72 (2H, t, J=7.6Hz), 6.42
(1H, d, J=3.5Hz), 7.13-7.30 (2H, m), 7.39 (1H, s), 7.88 (1H,
brs), 8.15 (1H, brs), 8.27 (1H, d, J=8.6Hz), 11.07 (1H, s)

Compound 1-294

¹H-NMR (DMSO-d₆) δ ppm: 2.00 (3H, s), 2.32 (3H, s), 7.09-7.72
15 (3H, m), 7.92-8.40 (3H, m), 10.95 (1H, s)

Compound 1-295

¹H-NMR (DMSO-d₆) δ ppm: 2.00 (3H, s), 2.32 (3H, s), 2.47 (3H,
s), 7.11-7.20 (2H, m), 7.38 (1H, s), 7.85 (1H, brs), 8.14 (1H,
brs), 8.26 (1H, d, J=8.1Hz), 10.98 (1H, s)

20 Compound 1-296

¹H-NMR (CDCl₃+MeOD-d₄) δ ppm: 4.67 (2H, s), 6.55-6.60 (1H, m),
7.15-7.30 (2H, m), 7.35-7.45 (1H, m), 8.25-8.35 (1H, m)

Compound 1-297

¹H-NMR (DMSO-d₆) δ ppm: 4.93 (2H, s), 6.81 (1H, d, J=3.5Hz),
25 7.34 (1H, d, J=3.5Hz), 7.43 (1H, dd, J=1.9, 9.1Hz), 7.75 (1H,
d, J=1.9Hz), 8.00 (1H, brs), 8.20-8.35 (2H, m), 11.07 (1H, s)

Compound 1-298

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 4.93 (2H, s), 6.82 (1H, d, $J=3.5\text{Hz}$), 7.36 (1H, d, $J=3.5\text{Hz}$), 7.55-7.65 (1H, m), 7.80-7.90 (1H, m), 8.08 (1H, brs), 8.36 (1H, brs), 11.00 (1H, s)

Compound 1-299

5 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 0.85-1.07 (4H, m), 2.04-2.15 (1H, m), 6.35-6.46 (1H, m), 7.18-7.28 (1H, m), 7.33-7.49 (1H, m), 7.57-7.70 (1H, m), 7.94-8.39 (3H, m), 11.08 (1H, s)

Compound 1-300

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 0.90-1.09 (4H, m), 2.05-2.17 (1H, m), 10 6.39-6.46 (1H, m), 7.20-7.33 (2H, m), 7.48-7.60 (1H, m), 8.00 (1H, brs), 8.24 (1H, brs), 8.43-8.50 (1H, m), 11.14 (1H, s)

Compound 1-301

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 0.90-1.09 (4H, m), 2.06-2.18 (1H, m), 6.38-6.47 (1H, m), 7.21-7.30 (1H, m), 7.66-7.76 (1H, m), 15 8.06-8.33 (3H, m), 11.03 (1H, s)

Compound 1-302

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 0.90-1.05 (4H, m), 2.05-2.15 (1H, m), 3.86 (3H, s), 6.42 (1H, d, $J=3.4\text{Hz}$), 6.95-7.10 (2H, m), 7.22 (1H, d, $J=3.4\text{Hz}$), 7.90 (1H, brs), 8.06 (1H, brs), 8.34 (1H, d, 20 $J=9.0\text{Hz}$), 11.19 (1H, s)

Compound 1-303

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.25 (6H, s), 2.47 (3H, s), 3.60 (2H, s), 6.55-6.65 (1H, m), 7.15-7.20 (1H, m), 7.25-7.45 (2H, m), 7.90 (1H, brs), 8.16 (1H, brs), 8.20-8.30 (1H, m), 11.08 (1H, 25 brs)

Compound 1-304

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.56 (3H, s), 7.35-7.70 (4H, m), 8.00-8.10

(1H, m), 8.11 (1H, brs), 8.35 (1H, brs), 11.38 (1H, brs)

Compound 1-305

¹H-NMR (DMSO-d₆) δ ppm: 1.40-1.75 (8H, m), 3.35-3.45 (4H, m),
7.30-7.75 (4H, m), 8.00-8.10 (1H, m), 8.10 (1H, brs), 8.35 (1H,
5 brs), 11.27 (1H, brs)

Compound 1-306

¹H-NMR (CDCl₃+MeOD-d₄) δ ppm: 2.32 (3H, s), 2.50-2.60 (4H, m),
3.35-3.45 (4H, m), 7.00-7.10 (1H, m), 7.15 (1H, d, J=3.7Hz),
7.36 (1H, d, J=3.7Hz), 8.05-8.15 (1H, m)

10 Compound 1-307

MS (ESI, m/z): 428 (M+H)+

Compound 1-308

MS (ESI, m/z): 476 (M+H)+

Compound 1-309

15 MS (ESI, m/z): 490 (M+H)+

Compound 1-310

¹H-NMR (DMSO-d₆) δ ppm: 7.35-7.70 (7H, m), 8.05-8.20 (2H, m),
8.13 (1H, brs), 8.36 (1H, brs), 11.39 (1H, brs)

Compound 1-311

20 ¹H-NMR (DMSO-d₆) δ ppm: 7.35-7.70 (6H, m), 7.85-7.95 (1H, m),
8.00-8.20 (3H, m), 8.35 (1H, brs), 11.39 (1H, brs)

Compound 1-312

¹H-NMR (DMSO-d₆) δ ppm: 7.05-7.25 (3H, m), 7.30-7.70 (3H, m),
7.90-8.00 (2H, m), 8.10-8.20 (2H, m), 8.36 (1H, brs), 11.40 (1H,
25 brs)

Compound 1-313

¹H-NMR (DMSO-d₆) δ ppm: 2.48 (3H, s), 7.15-7.75 (7H, m), 7.85-8.45

(4H, m), 11.45 (1H, brs)

Compound 1-314

¹H-NMR (DMSO-d₆) δ ppm: 2.48 (3H, s), 7.15-7.25 (1H, m), 7.35-7.60
 (5H, m), 7.85-8.05 (3H, m), 8.24 (1H, brs), 8.25-8.35 (1H, m),
 5 11.45 (1H, brs)

Compound 1-315

¹H-NMR (DMSO-d₆) δ ppm: 2.48 (3H, s), 7.15-7.25 (1H, m), 7.30-7.35
 (1H, m), 7.40-7.50 (2H, m), 7.55-7.65 (2H, m), 7.90-8.00 (2H,
 m), 8.03 (1H, brs), 8.25 (1H, brs), 8.25-8.35 (1H, m), 11.46
 10 (1H, brs)

Compound 1-316

¹H-NMR (DMSO-d₆) δ ppm: 4.83 (2H, s), 7.30 (1H, d, J=4.0Hz),
 7.45-7.70 (4H, m), 7.91 (1H, d, J=7.6Hz), 7.95-8.05 (2H, m),
 8.17 (1H, brs), 8.40 (1H, brs), 11.31 (1H, s)
 15 MS (ESI, m/z): 431 (M+H)+

Compound 1-317

¹H-NMR (DMSO-d₆) δ ppm: 2.48 (3H, s), 3.97 (3H, s), 7.05-7.25
 (4H, m), 7.35-7.50 (3H, m), 8.00-8.10 (2H, m), 8.34 (1H, brs),
 8.30-8.40 (1H, m), 11.52 (1H, brs)

20 Compound 1-318

¹H-NMR (DMSO-d₆) δ ppm: 2.48 (3H, s), 3.84 (3H, s), 7.00-7.25
 (4H, m), 7.35-7.45 (2H, m), 7.80-7.95 (2H, m), 8.02 (1H, brs),
 8.24 (1H, brs), 8.30-8.40 (1H, m), 11.48 (1H, brs)

Compound 1-319

25 ¹H-NMR (DMSO-d₆) δ ppm: 3.98 (3H, s), 7.05-7.25 (3H, m), 7.35-7.50
 (3H, m), 7.60-7.70 (1H, m), 8.00-8.10 (1H, m), 8.17 (1H, brs),
 8.20-8.25 (1H, m), 8.37 (1H, brs), 11.47 (1H, brs)

Compound 1-320

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.87 (3H, s), 6.95-7.05 (1H, m), 7.25-7.70 (7H, m), 8.14 (1H, brs), 8.20-8.30 (1H, m), 8.37 (1H, brs), 11.51 (1H, brs)

5 Compound 1-321

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.84 (3H, s), 7.05-7.15 (3H, m), 7.40-7.50 (1H, m), 7.60-7.70 (1H, m), 7.80-7.90 (3H, m), 8.14 (1H, brs), 8.20-8.25 (1H, m), 8.36 (1H, brs), 11.42 (1H, brs)

Compound 1-322

10 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.30-7.70 (5H, m), 7.90-8.05 (2H, m), 8.10-8.20 (1H, m), 8.12 (1H, brs), 8.34 (1H, brs), 8.65-8.70 (1H, m), 11.43 (1H, brs)

Compound 1-323

15 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.35-7.70 (5H, m), 8.15 (1H, brs), 8.15-8.35 (1H, m), 8.35 (1H, brs), 8.55-8.65 (1H, m), 9.15-9.25 (1H, m), 11.52 (1H, brs)

Compound 1-324

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.35-7.90 (7H, m), 8.10-8.20 (1H, m), 8.35 (1H, brs), 8.65-8.75 (2H, m), 11.51 (1H, brs)

20 Compound 1-325

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.48 (3H, s), 7.15-7.25 (1H, m), 7.35-7.60 (3H, m), 7.80-7.90 (2H, m), 8.05 (1H, brs), 8.24 (1H, brs), 8.25-8.40 (1H, m), 8.65-8.75 (2H, m), 11.58 (1H, brs)

Compound 1-326

25 MS (ESI, m/z): 382 (M-H) -

Compound 1-327

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.00-3.13 (4H, m), 6.39 (1H, d, J=3.5Hz),

7.14-7.70 (8H, m), 8.00-8.16 (2H, m), 8.32 (1H, brs), 11.13 (1H, s)

Compound 1-328

¹H-NMR (DMSO-d₆) δ ppm: 2.47 (3H, s), 3.00-3.10 (4H, m), 6.38
5 (1H, d, J=3.5Hz), 7.13-7.33 (7H, m), 7.40 (1H, s), 7.93 (1H, brs), 8.14-8.32 (2H, m), 11.17 (1H, s)

Compound 1-329

¹H-NMR (DMSO-d₆) δ ppm: 2.47 (3H, s), 2.99 (4H, s), 3.80 (3H, s), 6.30-6.44 (1H, m), 6.79-7.02 (2H, m), 7.10-7.45 (5H, m),
10 7.80-8.35 (3H, m), 11.12 (1H, s)

Compound 1-330

¹H-NMR (DMSO-d₆) δ ppm: 2.47 (3H, s), 2.95-3.15 (4H, m), 3.71 (3H, s), 6.33-6.44 (1H, m), 6.66-6.90 (3H, m), 7.09-7.45 (4H, m), 7.80-8.35 (3H, m), 11.16 (1H, s)

15 Compound 1-331

¹H-NMR (DMSO-d₆) δ ppm: 2.47 (3H, s), 2.90-3.10 (4H, m), 3.70 (3H, s), 6.30-6.45 (1H, m), 6.73-7.45 (7H, m), 7.80-8.35 (3H, m), 11.15 (1H, s)

Compound 1-332

20 ¹H-NMR (DMSO-d₆) δ ppm: 2.99 (4H, s), 3.81 (3H, s), 6.33-6.42 (1H, m), 6.80-7.03 (2H, m), 7.10-7.70 (5H, m), 8.01 (1H, brs), 8.07-8.16 (1H, m), 8.27 (1H, brs), 11.07 (1H, s)

Compound 1-333

¹H-NMR (DMSO-d₆) δ ppm: 2.90-3.15 (4H, m), 3.71 (3H, s), 6.39
25 (1H, d, J=3.5Hz), 6.70-6.90 (3H, m), 7.10-7.73 (4H, m), 8.03 (1H, brs), 8.08-8.17 (1H, m), 8.28 (1H, brs), 11.11 (1H, s)

Compound 1-334

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.90-3.10 (4H, m), 3.70 (3H, s), 6.37 (1H, d, $J=3.3\text{Hz}$), 6.75-6.90 (2H, m), 7.10-7.73 (5H, m), 8.03 (1H, brs), 8.08-8.17 (1H, m), 8.28 (1H, brs), 11.10 (1H, s)

Compound 1-335

5 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.40-7.50 (1H, m), 7.70-7.80 (2H, m), 7.95 (1H, brs), 8.10-8.30 (3H, m), 8.50-8.60 (1H, m), 8.75-8.80 (1H, m), 12.09 (1H, brs)

Compound 1-336

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 7.50-7.80 (2H, m), 8.03 (1H, brs),
10 8.05-8.30 (3H, m), 8.35 (1H, brs), 8.70-8.85 (1H, m), 12.05 (1H, brs)

Compound 1-337

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.87 (3H, s), 6.95-7.05 (1H, m), 7.09 (1H, d, $J=2.3\text{Hz}$), 7.60-7.90 (2H, m), 8.00 (1H, brs), 8.05-8.20
15 (1H, m), 8.24 (1H, d, $J=8.0\text{Hz}$), 8.46 (1H, d, $J=8.7\text{Hz}$), 8.77 (1H, d, $J=4.3\text{Hz}$), 12.11 (1H, s)

Compound 1-338

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 4.00 (3H, s), 7.10-7.20 (1H, m), 7.70-7.75 (1H, m), 7.75-7.85 (1H, m), 7.91 (1H, brs), 8.05-8.15 (1H, m),
20 8.15-8.30 (2H, m), 8.75-8.80 (1H, m), 12.03 (1H, brs)

Compound 1-339

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.62 (3H, s), 7.55-7.65 (2H, m), 7.95-8.10 (3H, m), 8.15-8.25 (1H, m), 8.35 (1H, brs), 12.26 (1H, brs)

Compound 1-340

25 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.30-1.40 (3H, m), 4.35-4.50 (2H, m), 6.95-7.05 (1H, m), 7.35-7.45 (1H, m), 7.60-7.75 (1H, m), 7.85-7.95 (1H, m), 8.00 (1H, brs), 8.15-8.35 (2H, m), 8.80-8.85

(1H, m), 10.90 (1H, s)

Compound 1-341

¹H-NMR (DMSO-d₆) δ ppm: 3.87 (3H, s), 6.99 (1H, dd, J=2.2, 8.8Hz),
7.11 (1H, d, J=2.2Hz), 7.15-7.35 (4H, m), 7.40-7.55 (2H, m),
5 7.88 (1H, brs), 8.07 (1H, brs), 8.12 (1H, d, J=8.8Hz), 8.38 (1H,
dd, J=2.5, 8.8Hz), 8.76 (1H, d, J=2.5Hz), 11.05 (1H, s)

Compound 1-342

¹H-NMR (DMSO-d₆) δ ppm: 1.43 (3H, t, J=6.9Hz), 4.29 (2H, q,
J=6.9Hz), 7.05-7.35 (6H, m), 7.40-7.65 (3H, m), 7.91 (1H, brs),
10 8.08 (1H, brs), 8.40 (1H, dd, J=2.5, 8.5Hz), 8.78 (1H, d, J=2.5Hz),
10.83 (1H, s)

Compound 1-343

¹H-NMR (DMSO-d₆) δ ppm: 2.48 (3H, s), 7.15-7.35 (5H, m), 7.38-7.55
(3H, m), 7.88 (1H, brs), 8.07 (1H, d, J=8.2Hz), 8.14 (1H, brs),
15 8.39 (1H, dd, J=2.5, 8.2Hz), 8.70-8.80 (1H, m), 10.97 (1H, s)

Compound 1-344

¹H-NMR (DMSO-d₆) δ ppm: 1.25 (3H, t, J=7.6Hz), 2.77 (2H, q,
J=7.6Hz), 7.15-7.35 (5H, m), 7.40-7.50 (3H, m), 7.89 (1H, brs),
8.07 (1H, d, J=8.5Hz), 8.12 (1H, brs), 8.39 (1H, dd, J=2.5, 8.5Hz),
20 8.77 (1H, d, J=2.5Hz), 10.96 (1H, s)

Compound 1-345

¹H-NMR (DMSO-d₆) δ ppm: 7.15-7.35 (4H, m), 7.40-7.55 (2H, m),
7.80-7.90 (1H, m), 7.99 (1H, brs), 8.10-8.20 (1H, m), 8.23 (1H,
brs), 8.35-8.45 (1H, m), 8.70-8.80 (1H, m), 10.95 (1H, brs)

25 Compound 1-346

¹H-NMR (DMSO-d₆) δ ppm: 7.15-7.35 (4H, m), 7.40-7.65 (3H, m),
7.70 (1H, dd, J=2.2, 8.8Hz), 8.07 (1H, brs), 8.33 (1H, brs),

8.40 (1H, dd, J=2.5, 8.8Hz), 8.77 (1H, d, J=2.5Hz), 10.87 (1H, s)

Compound 1-347

¹H-NMR (DMSO-d₆) δ ppm: 2.39 (3H, s), 7.15-7.35 (4H, m), 7.40-7.60
5 (3H, m), 7.87 (1H, d, J=10.1Hz), 7.97 (1H, brs), 8.23 (1H, brs),
8.39 (1H, dd, J=2.2, 8.5Hz), 8.76 (1H, d, J=2.2Hz), 10.97 (1H, s)

Compound 1-348

¹H-NMR (DMSO-d₆) δ ppm: 4.00 (3H, s), 7.05-7.55 (8H, m), 7.95
10 (1H, brs), 8.17 (1H, brs), 8.35-8.45 (1H, m), 8.70-8.80 (1H, m),
10.88 (1H, brs)

Compound 1-349

¹H-NMR (DMSO-d₆) δ ppm: 7.25-7.50 (6H, m), 7.60-7.75 (1H, m),
7.86 (1H, dd, J=2.8, 9.1Hz), 7.97 (1H, brs), 8.22 (1H, brs),
15 8.43 (1H, dd, J=2.5, 8.5Hz), 8.74 (1H, d, J=2.5Hz), 10.88 (1H, s)

Compound 1-350

¹H-NMR (DMSO-d₆) δ ppm: 7.00-7.33 (4H, m), 7.35-7.55 (2H, m),
7.63-7.73 (1H, m), 7.87 (1H, dd, J=2.8, 9.1Hz), 7.98 (1H, brs),
20 8.24 (1H, brs), 8.42 (1H, brs), 8.79 (1H, d, J=2.2Hz), 10.91
(1H, s)

Compound 1-351

¹H-NMR (DMSO-d₆) δ ppm: 7.15-7.45 (6H, m), 7.60-7.70 (1H, m),
7.87 (1H, dd, J=2.5, 9.5Hz), 7.98 (1H, brs), 8.23 (1H, brs),
25 8.35-8.45 (1H, m), 8.76 (1H, d, J=2.5Hz), 10.90 (1H, s)

Compound 1-352

¹H-NMR (DMSO-d₆) δ ppm: 7.10-7.25 (1H, m), 7.30-7.55 (4H, m),

7.67 (1H, dd, J=4.1, 8.8Hz), 7.85 (1H, dd, J=2.5, 9.1Hz), 7.98 (1H, brs), 8.25 (1H, brs), 8.43 (1H, dd, J=2.2, 8.8 Hz), 8.74 (1H, dd, J=2.2 Hz), 10.88 (1H, s)

Compound 1-353

- 5 $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 7.30-7.55 (4H, m), 7.68 (1H, dd, J=4.1, 9.1 Hz), 7.83 (1H, dd, J=2.5, 9.1Hz), 7.98 (1H, brs), 8.24 (1H, brs), 8.48 (1H, dd, J=2.2, 8.5 Hz), 8.74 (1H, d, J=2.2Hz), 10.88 (1H, s)

Compound 1-354

- 10 $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 1.15 (6H, d, J=6.9Hz), 2.95-3.10 (1H, m), 7.05-7.13 (1H, m), 7.15-7.33 (3H, m), 7.35-7.50 (2H, m), 7.67 (1H, dd, J=4.1, 9.1 Hz), 7.86 (1H, dd, J=2.5, 9.1 Hz), 7.97 (1H, brs), 8.22 (1H, brs), 8.39 (1H, dd, J=2.5, 8.5Hz), 8.75 (1H, d, J=2.5Hz), 10.87 (1H, s)

- 15 Compound 1-355

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 1.24 (6H, d, J=6.9Hz), 2.85-3.05 (1H, m), 7.05-7.25 (3H, m), 7.30-7.45 (3H, m), 7.60-7.70 (1H, m), 7.88 (1H, dd, J=2.8, 9.1Hz), 7.98 (1H, brs), 8.24 (1H, brs), 8.38 (1H, dd, J=2.5, 8.8Hz), 8.76 (1H, d, J=2.5Hz), 10.90 (1H, s)

- 20 Compound 1-356

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 2.10 (3H, s), 7.05-7.30 (4H, m), 7.35-7.45 (1H, m), 7.67 (1H, dd, J=4.1, 9.1Hz), 7.86 (1H, dd, J=2.8, 9.1Hz), 7.99 (1H, brs), 8.25 (1H, brs), 8.40 (1H, dd, J=2.5, 8.8Hz), 8.73 (1H, d, J=2.5Hz), 10.88 (1H, s)

- 25 Compound 1-357

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 4.80 (2H, s), 7.15-7.75 (7H, m), 8.04 (1H, brs), 8.30 (1H, brs), 8.35-8.45 (1H, m), 8.75-8.85 (1H,

m), 10.86 (1H, brs)

Compound 1-358

¹H-NMR (DMSO-d₆) δ ppm: 3.71 (3H, s), 6.95-7.45 (6H, m), 7.67
(1H, dd, J=4.1, 9.1Hz), 7.87 (1H, dd, J=2.8, 9.1Hz), 7.97 (1H,
5 brs), 8.23 (1H, brs), 8.36 (1H, dd, J=2.2, 8.5Hz), 8.71 (1H,
d, J=2.2Hz), 10.87 (1H, s)

Compound 1-359

¹H-NMR (DMSO-d₆) δ ppm: 3.77 (3H, s), 6.70-7.00 (3H, m), 7.20
(1H, d, J=8.5Hz), 7.30-7.50 (2H, m), 7.60-7.75 (1H, m), 7.80-7.92
10 (1H, m), 7.98 (1H, brs), 8.23 (1H, brs), 8.39 (1H, dd, J=1.9,
8.5Hz), 8.79 (1H, s), 10.90 (1H, s)

Compound 1-360

¹H-NMR (DMSO-d₆) δ ppm: 3.79 (3H, s), 6.95-7.05 (2H, m), 7.10-7.20
(3H, m), 7.35-7.45 (1H, m), 7.63-7.70 (1H, m), 7.85-7.93 (1H,
15 m), 7.98 (1H, brs), 8.23 (1H, brs), 8.37 (1H, dd, J=2.5, 8.8
Hz), 8.76 (1H, d, J=2.5 Hz), 10.89 (1H, s)

Compound 1-361

¹H-NMR (DMSO-d₆) δ ppm: 3.73 (3H, s), 6.75-6.90 (1H, m), 7.05-7.30
(3H, m), 7.35-7.45 (1H, m), 7.67 (1H, dd, J=4.1, 9.1Hz), 7.87
20 (1H, dd, J=2.5, 9.1Hz), 7.97 (1H, brs), 8.23 (1H, brs), 8.36
(1H, dd, J=2.5, 8.8Hz), 8.71 (1H, d, J=2.5Hz), 10.86 (1H, s)

Compound 1-362

¹H-NMR (DMSO-d₆) δ ppm: 7.30-7.47 (2H, m), 7.48-7.60 (1H, m),
7.60-7.80 (2H, m), 7.86 (1H, dd, J=2.5, 9.1Hz), 7.98 (1H, brs),
25 8.24 (1H, brs), 8.44 (1H, dd, J=2.5, 8.5Hz), 8.47-8.58 (2H, m),
8.76 (1H, d, J=2.2Hz), 10.91 (1H, s)

Compound 1-363

¹H-NMR (DMSO-d₆) δ ppm: 2.16 (3H, s), 7.45-7.60 (2H, m), 7.93 (1H, brs), 8.14 (1H, brs), 10.08 (1H, brs)

Compound 1-364

¹H-NMR (DMSO-d₆) δ ppm: 1.13 (3H, t, J=7.5Hz), 2.46 (2H, q, J=7.5Hz), 7.45-7.65 (2H, m), 7.92 (1H, brs), 8.19 (1H, brs), 10.09 (1H, s)

Compound 1-365

¹H-NMR (DMSO-d₆) δ ppm: 1.18 (6H, d, J=7.0Hz), 2.65-2.80 (1H, m), 7.45-7.70 (2H, m), 7.95 (1H, brs), 8.19 (1H, brs), 10.15 (1H, s)

MS (ESI, m/z): 283 (M+H)⁺

Compound 1-366

¹H-NMR (DMSO-d₆) δ ppm: 1.28 (9H, s), 7.50-7.60 (1H, m), 7.80-7.90 (1H, m), 8.03 (1H, brs), 8.29 (1H, brs), 10.43 (1H, s)

15 Compound 1-367

¹H-NMR (DMSO-d₆) δ ppm: 4.29 (2H, s), 7.45-7.62 (2H, m), 8.02 (1H, brs), 8.25 (1H, brs), 10.65 (1H, s)

Compound 1-368

¹H-NMR (DMSO-d₆) δ ppm: 1.70-1.80 (2H, m), 1.85-1.95 (2H, m), 2.40-2.60 (2H, m), 3.59 (2H, t, J=6.7Hz), 7.45-7.60 (2H, m), 7.94 (1H, brs), 8.16 (1H, brs), 10.10 (1H, s)

Compound 1-369

¹H-NMR (DMSO-d₆) δ ppm: 1.40-1.55 (2H, m), 1.60-1.75 (2H, m), 1.80-1.95 (2H, m), 2.46 (2H, t, J=7.4Hz), 3.55 (2H, t, J=6.8Hz), 7.45-7.60 (2H, m), 7.93 (1H, brs), 8.15 (1H, brs), 10.09 (1H, s)

Compound 1-370

¹H-NMR (DMSO-d₆) δ ppm: 3.08 (2H, t, J=7.0Hz), 4.59 (2H, t, J=7.0Hz), 6.90-7.05 (1H, m), 7.10-7.75 (7H, m), 7.80-8.35 (4H, m), 8.75-8.90 (1H, m), 10.90 (1H, s)

5 Example 3

6-Methoxy-3-[3-(4-phenylpiperazin-1-ylmethyl)benzoylamino]-benzofuran-2-carboxamide (compound 2-1)

Triethylamine (0.035mL) and 4-phenylpiperazine (76mg) were added to a solution of 3-(3-chloromethylbenzoylamino)-
10 6-methoxybenzofuran-2-carboxamide (compound 1-57, 73mg) in tetrahydrofuran (1mL), and the resulting mixture was stirred at 60°C overnight. Water was added to the reaction mixture, and the resulting precipitate was collected by filtration to give the title compound (67mg).

15 ¹H-NMR (DMSO-d₆) δ ppm: 2.50-2.60 (4H, m), 3.10-3.20 (4H, m), 3.30 (2H, s), 3.87 (3H, s), 6.70-6.80 (1H, m), 6.85-7.15 (6H, m), 7.55-7.70 (2H, m), 7.75-8.15 (4H, m), 8.20-8.30 (1H, m), 11.19 (1H, brs)

20 Compounds 2-2 to 2-145 were prepared in a manner similar to those as described in Example 3 using the corresponding benzylhalides or 2-chloromethylfurans, and amines or thiols instead of 3-(3-chloromethylbenzoylamino)-6-methoxy-benzofuran-2-carboxamide (compound 1-57) and
25 4-phenylpiperazine. These were illustrated in table 3.

Table 3

Compound No.	Structure	Compound No.	Structure
2-1		2-9	
2-2		2-10	
2-3		2-11	
2-4		2-12	
2-5		2-13	
2-6		2-14	
2-7		2-15	
2-8		2-16	

Table 3 (continued)

Compound No.	Structure	Compound No.	Structure
2-17		2-25	
2-18		2-26	
2-19		2-27	
2-20		2-28	
2-21		2-29	
2-22		2-30	
2-23		2-31	
2-24		2-32	

Table 3 (continued)

Compound No.	Structure	Compound No.	Structure
2-33		2-41	
2-34		2-42	
2-35		2-43	
2-36		2-44	
2-37		2-45	
2-38		2-46	
2-39		2-47	
2-40		2-48	

Table 3 (continued)

Compound No.	Structure	Compound No.	Structure
2-49		2-57	
2-50		2-58	
2-51		2-59	
2-52		2-60	
2-53		2-61	
2-54		2-62	
2-55		2-63	
2-56		2-64	

Table 3 (continued)

Compound No.	Structure	Compound No.	Structure
2-65		2-73	
2-66		2-74	
2-67		2-75	
2-68		2-76	
2-69		2-77	
2-70		2-78	
2-71		2-79	
2-72		2-80	

Table 3 (continued)

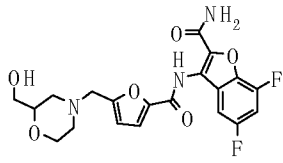
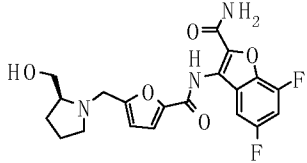
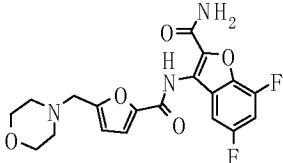
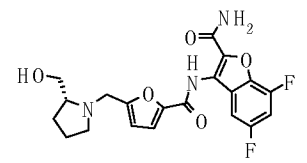
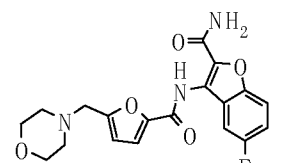
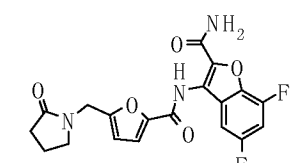
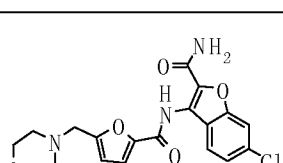
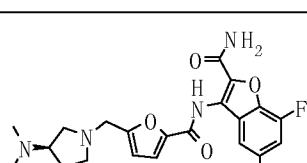
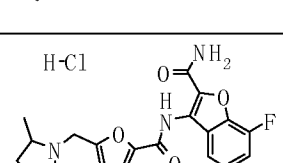
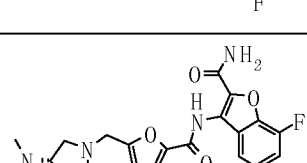
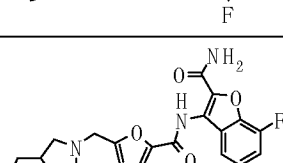
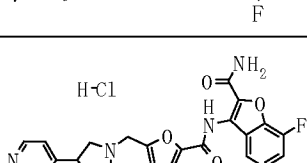
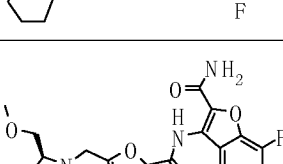
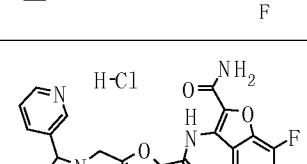
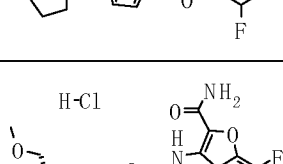
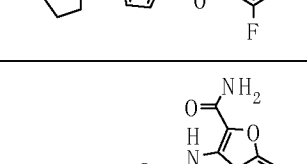
Compound No.	Structure	Compound No.	Structure
2-81		2-89	
2-82		2-90	
2-83		2-91	
2-84		2-92	
2-85	 H·Cl	2-93	
2-86		2-94	 H·Cl
2-87		2-95	 H·Cl
2-88	 H·Cl	2-96	

Table 3 (continued)

Compound No.	Structure	Compound No.	Structure
2-97		2-105	
2-98		2-106	
2-99		2-107	
2-100		2-108	
2-101		2-109	
2-102		2-110	
2-103		2-111	
2-104		2-112	

Table 3 (continued)

Compound No.	Structure	Compound No.	Structure
2-113		2-121	
2-114		2-122	
2-115		2-123	
2-116		2-124	
2-117		2-125	
2-118		2-126	
2-119		2-127	
2-120		2-128	

Table 3 (continued)

Compound No.	Structure	Compound No.	Structure
2-129		2-138	
2-130		2-139	
2-131		2-140	
2-132		2-141	
2-133		2-142	
2-134		2-143	
2-135		2-144	
2-136		2-145	
2-137			

The physical data of compounds 2-2 to 2-145 were described below.

Compound 2-2

¹H-NMR (DMSO-d₆) δ ppm: 1.65-1.75 (4H, m), 2.40-2.50 (4H, m),
5 3.68 (2H, s), 3.87 (3H, s), 6.95-7.15 (2H, m), 7.50-7.65 (2H, m), 7.80-7.95 (3H, m), 8.06 (1H, brs), 8.20-8.30 (1H, m), 11.18 (1H, brs)

Compound 2-3

¹H-NMR (DMSO-d₆) δ ppm: 0.85 (6H, d, J=6.6Hz), 1.30-1.40 (2H,
10 m), 1.64 (1H, heptet, J=6.6Hz), 2.08 (1H, brs), 2.45-2.55 (2H, m), 3.78 (2H, s), 3.87 (3H, s), 6.95-7.15 (2H, m), 7.50-7.65 (2H, m), 7.80-8.15 (4H, m), 8.20-8.30 (1H, m), 11.14 (1H, brs)

Compound 2-4

¹H-NMR (DMSO-d₆) δ ppm: 1.00-1.25 (5H, m), 1.50-2.05 (6H, m),
15 2.30-2.45 (1H, m), 3.82 (2H, s), 3.87 (3H, s), 6.95-7.15 (2H, m), 7.45-7.70 (2H, m), 7.80-8.10 (3H, m), 8.05 (1H, brs), 8.20-8.30 (1H, m), 11.14 (1H, brs)

Compound 2-5

¹H-NMR (DMSO-d₆) δ ppm: 3.86 (3H, s), 4.30 (2H, s), 6.90-7.25
20 (4H, m), 7.35-7.55 (4H, m), 7.80-7.95 (3H, m), 8.04 (1H, brs), 8.15-8.25 (1H, m), 11.11 (1H, brs)

Compound 2-6

¹H-NMR (DMSO-d₆) δ ppm: 3.87 (3H, s), 4.32 (2H, s), 6.90-7.20
25 (4H, m), 7.40-7.50 (2H, m), 7.80-7.95 (3H, m), 8.05 (1H, brs), 8.20-8.30 (1H, m), 11.11 (1H, brs), 12.25 (1H, brs)

Compound 2-7

¹H-NMR (DMSO-d₆) δ ppm: 3.39 (3H, s), 3.87 (3H, s), 4.27 (2H,

s), 6.95-7.15 (3H, m), 7.20-7.25 (1H, m), 7.35-7.45 (2H, m), 7.80-7.90 (3H, m), 8.05 (1H, brs), 8.20-8.30 (1H, m), 11.13 (1H, brs)

Compound 2-8

- 5 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.55-2.70 (4H, m), 3.05-3.25 (4H, m), 3.70 (2H, s), 3.86 (3H, s), 6.60-6.65 (1H, m), 6.70-6.80 (1H, m), 6.85-7.25 (6H, m), 7.30-7.35 (1H, m), 7.88 (1H, brs), 8.08 (1H, brs), 8.25-8.30 (1H, m), 11.13 (1H, brs)

Compound 2-9

- 10 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.10-2.25 (6H, m), 2.65-2.75 (2H, m), 3.80-3.95 (5H, m), 6.55-6.65 (1H, m), 6.95-7.15 (2H, m), 7.25-7.40 (1H, m), 7.86 (1H, brs), 8.08 (1H, brs), 8.20-8.30 (1H, m), 11.11 (1H, brs)

Compound 2-10

- 15 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.86 (3H, s), 4.35 (2H, s), 6.40-6.50 (1H, m), 6.90-7.25 (5H, m), 7.40-7.55 (2H, m), 7.89 (1H, brs), 8.08 (1H, brs), 8.20-8.30 (1H, m), 11.08 (1H, brs)

Compound 2-11

- 20 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 3.53 (3H, s), 4.31 (2H, s), 6.35-6.40 (1H, m), 6.95-7.00 (1H, m), 7.20-7.30 (2H, m), 7.35-7.45 (1H, m), 7.60-7.70 (1H, m), 7.95-8.10 (1H, m), 8.01 (1H, brs), 8.27 (1H, brs), 10.95 (1H, brs)

Compound 2-12

- 25 $^1\text{H-NMR}$ (CDCl_3) δ ppm: 2.28 (6H, s), 3.53 (2H, s), 5.65 (1H, brs), 6.40 (1H, brs), 7.25-7.40 (1H, m), 7.45-7.65 (3H, m), 7.90-8.00 (2H, m), 8.55-8.65 (1H, m), 10.88 (1H, brs)

Compound 2-13

¹H-NMR (DMSO-d₆) δ ppm: 2.19 (6H, s), 3.51 (2H, s), 7.35-7.50 (1H, m), 7.50-7.75 (3H, m), 7.85-7.95 (1H, m), 7.95 (1H, s), 7.95-8.10 (2H, m), 8.31 (1H, brs), 11.05 (1H, s)

MS (ESI, m/z): 356 (M+H)+

5 Compound 2-14

MS (ESI, m/z): 470 (M+H)+

Compound 2-15

¹H-NMR (DMSO-d₆) δ ppm: 2.22 (3H, s), 2.70-2.80 (2H, m), 2.90-3.00 (2H, m), 3.63 (2H, s), 7.10-7.20 (1H, m), 7.28 (1H, d, J=7.6Hz),
10 7.45-7.70 (4H, m), 7.80-7.95 (3H, m), 8.09 (1H, brs), 8.36 (1H, brs), 8.44 (1H, dd, J=0.9, 4.7Hz), 10.96 (1H, s)

Compound 2-16

¹H-NMR (DMSO-d₆) δ ppm: 2.20 (3H, s), 2.50-2.65 (2H, m), 3.24 (3H, s), 3.42-3.70 (4H, m), 7.48-7.70 (3H, m), 7.76-8.00 (3H, m),
15 8.09 (1H, s), 8.36 (1H, s), 10.98 (1H, s)

Compound 2-17

MS (ESI, m/z): 404 (M+H)+

Compound 2-18

¹H-NMR (CDCl₃) δ ppm: 1.16 (9H, s), 2.82 (2H, t, J=5.5Hz), 3.63 (2H, t, J=5.5Hz), 3.82 (2H, s), 6.68 (1H, brs), 6.95-7.10 (1H, m),
20 7.40-7.60 (2H, m), 7.85-7.95 (1H, m), 8.10-8.30 (2H, m), 10.99 (1H, brs)

Compound 2-19

¹H-NMR (DMSO-d₆) δ ppm: 0.90-1.35 (5H, m), 1.45-1.90 (7H, m),
25 2.30-2.70 (3H, m), 3.73 (2H, s), 4.20-4.35 (1H, m), 7.45-7.70 (3H, m), 7.80-7.90 (2H, m), 7.97 (1H, s), 8.10 (1H, brs), 8.40 (1H, brs), 11.00 (1H, brs)

Compound 2-20

$^1\text{H-NMR}$ ($\text{CDCl}_3 + \text{MeOD-d}_4$) δ ppm: 2.65-2.75 (4H, m), 3.60-3.70 (4H, m), 3.78 (2H, s), 7.00-7.10 (1H, m), 7.45-7.55 (2H, m), 7.90-8.00 (1H, m), 8.15-8.25 (1H, m)

5 Compound 2-21

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.67 (4H, t, $J=5.9\text{Hz}$), 3.21 (6H, s), 3.43 (4H, t, $J=5.9\text{Hz}$), 3.76 (2H, s), 7.50-7.65 (3H, m), 7.80-7.95 (2H, m), 7.96 (1H, s), 8.10 (1H, brs), 8.38 (1H, brs), 10.98 (1H, s)

10 MS (ESI, m/z): 462 (M+H)+

Compound 2-22

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.30-2.45 (4H, m), 3.50-3.65 (6H, m), 7.35-7.45 (1H, m), 7.50-7.70 (3H, m), 7.85-8.10 (4H, m), 8.27 (1H, brs), 11.04 (1H, s)

15 MS (ESI, m/z): 398 (M+H)+

Compound 2-23

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.30-2.45 (4H, m), 3.50-3.65 (6H, m), 7.40-7.50 (1H, m), 7.50-7.65 (2H, m), 7.70-7.80 (1H, m), 7.90 (1H, d, $J=7.8\text{Hz}$), 7.90-8.10 (2H, m), 8.20-8.35 (2H, m), 11.07

20 (1H, s)

MS (ESI, m/z): 414 (M+H)+

Compound 2-24

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.40 (4H, brs), 3.50-3.70 (6H, m), 7.50-7.70 (3H, m), 7.80-7.90 (1H, m), 7.90 (1H, d, $J=7.5\text{Hz}$), 7.96 (1H, s), 8.10 (1H, brs), 8.38 (1H, brs), 11.00 (1H, s)

25

MS (ESI, m/z): 416 (M+H)+

Compound 2-25

¹H-NMR (DMSO-d₆) δ ppm: 1.82 (1H, t, J=10.5Hz), 2.00-2.15 (1H, m), 2.64 (1H, d, J=11.0Hz), 2.78 (1H, d, J=11.0Hz), 3.20-3.65 (6H, m), 3.77 (1H, d, J=11.1Hz), 4.65 (1H, t, J=6.0Hz), 7.50-7.65 (3H, m), 7.83 (1H, d, J=7.8Hz), 7.91 (1H, d, J=7.3Hz), 7.96 (1H, s), 8.11 (1H, brs), 8.38 (1H, brs), 10.99 (1H, s)

MS (ESI, m/z): 446 (M+H)+

Compound 2-26

¹H-NMR (DMSO-d₆) δ ppm: 1.13 (3H, d, J=6.0Hz), 1.30-1.45 (1H, m), 1.50-1.75 (2H, m), 1.85-2.20 (2H, m), 2.35-2.50 (1H, m), 2.75-2.85 (1H, m), 3.23 (1H, d, J=13.2Hz), 4.07 (1H, d, J=13.2Hz), 7.45-7.65 (3H, m), 7.80-8.00 (3H, m), 8.10 (1H, brs), 8.37 (1H, brs), 11.00 (1H, s)

Compound 2-27

¹H-NMR (DMSO-d₆) δ ppm: 1.90-2.05 (2H, m), 2.34 (2H, t, J=8.1Hz), 3.29 (2H, t, J=7.1Hz), 4.49 (2H, s), 7.52 (1H, d, J=7.8Hz), 7.55-7.65 (2H, m), 7.75-7.90 (2H, m), 7.92 (1H, d, J=7.8Hz), 8.10 (1H, brs), 8.38 (1H, brs), 11.00 (1H, s)

MS (ESI, m/z): 414 (M+H)+

Compound 2-28

¹H-NMR (DMSO-d₆) δ ppm: 2.74 (4H, s), 4.66 (2H, s), 7.50-7.70 (3H, m), 7.80-7.95 (3H, m), 8.11 (1H, brs), 8.39 (1H, brs), 11.00 (1H, brs)

Compound 2-29

MS (ESI, m/z): 443 (M+H)+

Compound 2-30

¹H-NMR (DMSO-d₆) δ ppm: 1.40-2.00 (4H, m), 2.10-2.30 (1H, m), 2.60-2.90 (2H, m), 3.20-3.50 (6H, m), 4.10-4.30 (1H, m),

7.50-7.65 (3H, m), 7.80-8.00 (3H, m), 8.09 (1H, brs), 8.37 (1H, brs), 10.99 (1H, s)

MS (ESI, m/z): 444 (M+H)+

Compound 2-31

5 $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 1.40-2.00 (4H, m), 2.10-2.30 (1H, m), 2.60-2.90 (2H, m), 3.20-3.50 (6H, m), 4.10-4.30 (1H, m), 7.50-7.65 (3H, m), 7.80-8.00 (3H, m), 8.09 (1H, brs), 8.37 (1H, brs), 10.99 (1H, s)

Compound 2-32

10 MS (ESI, m/z): 430 (M+H)+

Compound 2-33

MS (ESI, m/z): 430 (M+H)+

Compound 2-34

15 $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 1.70-1.90 (1H, m), 2.20-3.05 (5H, m), 3.25-3.45 (1H, m), 3.77 (2H, brs), 7.25-7.40 (2H, m), 7.50-7.70 (3H, m), 7.80-7.95 (2H, m), 8.00 (1H, s), 8.10 (1H, brs), 8.39 (1H, brs), 8.42-8.50 (2H, m), 11.01 (1H, s)

Compound 2-35

20 $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 1.30-1.45 (2H, m), 1.45-1.60 (4H, m), 2.35 (4H, brs), 3.53 (2H, s), 7.35-7.45 (1H, m), 7.50-7.75 (3H, m), 7.80-8.10 (4H, m), 8.27 (1H, brs), 11.03 (1H, s)

MS (ESI, m/z): 396 (M+H)+

Compound 2-36

25 $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 1.30-1.60 (6H, m), 2.36 (4H, brs), 3.54 (2H, s), 7.45-7.65 (3H, m), 7.75-8.00 (3H, m), 8.09 (1H, brs), 8.37 (1H, brs), 10.99 (1H, brs)

MS (ESI, m/z): 414 (M+H)+

Compound 2-37

¹H-NMR (DMSO-d₆) δ ppm: 1.12 (3H, d, J=6.0Hz), 1.15-1.70 (6H, m), 1.90-2.05 (1H, m), 2.25-2.45 (1H, m), 2.55-2.70 (1H, m), 3.25 (1H, d, J=14Hz), 4.01 (1H, d, J=14Hz), 7.50-7.65 (3H, m),
5 7.80-7.90 (2H, m), 7.94 (1H, s), 8.11 (1H, brs), 8.38 (1H, brs), 11.00 (1H, brs)

MS (ESI, m/z): 428 (M+H)⁺

Compound 2-38

¹H-NMR (DMSO-d₆) δ ppm: 0.88 (3H, t, J=7.6Hz), 1.00-1.80 (8H, m),
10 1.90-2.80 (3H, m), 3.20-3.40 (1H, m), 3.90-4.10 (1H, m), 7.45-7.70 (3H, m), 7.80-8.00 (3H, m), 8.09 (1H, brs), 8.36 (1H, brs), 10.99 (1H, s)

Compound 2-39

¹H-NMR (DMSO-d₆) δ ppm: 1.30-1.50 (2H, m), 1.65-1.80 (2H, m),
15 2.00-2.15 (2H, m), 2.60-2.75 (2H, m), 3.40-3.60 (3H, m), 4.50-4.60 (1H, m), 7.50-7.70 (3H, m), 7.80-8.00 (3H, m), 8.10 (1H, brs), 8.38 (1H, brs), 10.98 (1H, s)

MS (ESI, m/z): 430 (M+H)⁺

Compound 2-40

¹H-NMR (CDCl₃+MeOD-d₄) δ ppm: 1.20-1.80 (5H, m), 1.90-2.15 (2H, m),
20 2.85-3.00 (2H, m), 3.30-3.50 (2H, m), 3.61 (2H, s), 7.00-7.10 (1H, m), 7.45-7.60 (2H, m), 7.85-8.00 (2H, m), 8.10-8.20 (1H, m)

Compound 2-41

¹H-NMR (DMSO-d₆) δ ppm: 0.80-2.00 (7H, m), 2.60-2.95 (2H, m),
25 3.10-3.60 (4H, m), 4.30-4.45 (1H, m), 7.50-7.65 (3H, m), 7.80-7.95 (3H, m), 8.08 (1H, brs), 8.36 (1H, brs), 10.98 (1H, s)

s)

Compound 2-42

MS (ESI, m/z): 444 (M+H)+

Compound 2-43

5 MS (ESI, m/z): 458 (M+H)+

Compound 2-44

MS (ESI, m/z): 457 (M+H)+

Compound 2-45

¹H-NMR (DMSO-d₆) δ ppm: 1.25-1.40 (1H, m), 1.40-1.55 (1H, m),
 10 1.55-1.80 (2H, m), 1.85-2.10 (2H, m), 2.25-2.40 (1H, m),
 2.65-2.85 (2H, m), 3.50-3.65 (2H, m), 6.76 (1H, brs), 7.28 (1H,
 brs), 7.50-7.65 (3H, m), 7.80-8.00 (3H, m), 8.10 (1H, brs), 8.39
 (1H, brs), 10.99 (1H, s)

MS (ESI, m/z): 457 (M+H)+

15 Compound 2-46

MS (ESI, m/z): 462 (M+H)+

Compound 2-47

¹H-NMR (DMSO-d₆) δ ppm: 1.50-1.70 (4H, m), 3.62 (2H, s), 3.80-3.90
 (4H, m), 7.35-7.50 (1H, m), 7.50-7.65 (2H, m), 7.70-7.80 (1H,
 20 m), 7.85-8.10 (3H, m), 8.15-8.40 (2H, m), 11.05 (1H, s)

MS (ESI, m/z): 470 (M+H)+

Compound 2-48

¹H-NMR (DMSO-d₆) δ ppm: 1.55-1.70 (4H, m), 2.35-2.60 (4H, m),
 3.59 (2H, s), 3.85 (4H, s), 7.50-7.70 (3H, m), 7.75-8.00 (3H,
 25 m), 8.10 (1H, brs), 8.38 (1H, brs), 10.97 (1H, brs)

MS (ESI, m/z): 472 (M+H)+

Compound 2-49

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.05-2.60 (10H, m), 3.56 (2H, s), 7.50-7.70 (3H, m), 7.80-8.00 (3H, m), 8.10 (1H, brs), 8.38 (1H, brs), 10.98 (1H, s)

MS (ESI, m/z): 429 (M+H)+

5 Compound 2-50

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.20-2.60 (8H, m), 3.47 (2H, s), 3.58 (2H, s), 7.20-7.40 (5H, m), 7.40-7.50 (1H, m), 7.50-7.65 (2H, m), 7.70-7.80 (1H, m), 7.85-8.15 (3H, m), 8.15-8.40 (2H, m), 11.05 (1H, s)

10 MS (ESI, m/z): 503 (M+H)+

Compound 2-51

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.20-2.60 (8H, m), 3.46 (2H, s), 3.57 (2H, s), 7.15-7.35 (5H, m), 7.50-7.65 (3H, m), 7.80-8.00 (3H, m), 8.07 (1H, brs), 8.34 (1H, brs), 10.97 (1H, s)

15 MS (ESI, m/z): 505 (M+H)+

Compound 2-52

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 0.85-1.10 (6H, m), 2.20-2.75 (9H, m), 3.55 (2H, s), 7.45-7.65 (3H, m), 7.75-8.00 (3H, m), 8.06 (1H, brs), 8.34 (1H, brs), 10.97 (1H, s)

20 MS (ESI, m/z): 457 (M+H)+

Compound 2-53

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 2.30-2.70 (8H, m), 2.56 (2H, t, $J=7.4\text{Hz}$), 2.74 (2H, t, $J=7.4\text{Hz}$), 3.60 (2H, s), 5.13 (2H, s), 5.96 (1H, brs), 6.56 (1H, brs), 7.00-7.10 (1H, m), 7.25-7.60 (7H, m), 7.85-8.05 (2H, m), 8.15-8.25 (1H, m), 10.82 (1H, brs)

25

Compound 2-54

MS (ESI, m/z): 487 (M+H)+

Compound 2-55

¹H-NMR (DMSO-d₆) δ ppm: 3.25-3.55 (2H, m), 3.60-3.75 (2H, m),
4.05-4.25 (2H, m), 4.66 (2H, s), 5.11 (2H, s), 7.25-7.45 (5H,
m), 7.50-7.70 (3H, m), 7.75-8.00 (3H, m), 8.10 (1H, brs), 8.38
5 (1H, brs), 11.01 (1H, s)

MS (ESI, m/z): 563 (M+H)+

Compound 2-56

¹H-NMR (DMSO-d₆) δ ppm: 1.01 (6H, d, J=6.0Hz), 2.70-2.85 (1H,
m), 3.79 (2H, s), 6.45-6.60 (1H, m), 7.20-7.35 (1H, m), 7.35-7.50
10 (1H, m), 7.60-7.70 (1H, m), 7.90-8.15 (2H, m), 8.29 (1H, brs),
10.97 (1H, s)

MS (ESI, m/z): 360 (M+H)+

Compound 2-57

¹H-NMR (DMSO-d₆) δ ppm: 1.09 (9H, s), 3.78 (2H, s), 6.45-6.60
15 (1H, m), 7.25-7.35 (1H, m), 7.35-7.50 (1H, m), 7.60-7.75 (1H,
m), 7.95-8.20 (2H, m), 8.29 (1H, brs), 10.96 (1H, s)

MS (ESI, m/z): 374 (M+H)+

Compound 2-58

¹H-NMR (DMSO-d₆) δ ppm: 1.50-1.70 (12H, m), 1.95-2.10 (3H, m),
20 3.80 (2H, s), 6.45-6.55 (1H, m), 7.25-7.35 (1H, m), 7.50-7.65
(1H, m), 7.85-7.95 (1H, m), 8.05 (1H, brs), 8.35 (1H, brs), 10.89
(1H, brs)

Compound 2-59

¹H-NMR (DMSO-d₆) δ ppm: 1.20-1.35 (2H, m), 1.75-2.05 (4H, m),
25 2.35-2.55 (1H, m), 2.65-2.80 (2H, m), 3.43 (2H, s), 3.82 (2H,
s), 6.54 (1H, d, J=3.1Hz), 7.20-7.35 (6H, m), 7.35-7.45 (1H,
m), 7.60-7.70 (1H, m), 7.95-8.10 (2H, m), 8.29 (1H, brs), 10.96

(1H, s)

MS (ESI, m/z): 491 (M+H)+

Compound 2-60

¹H-NMR (DMSO-d₆) δ ppm: 2.23 (6H, s), 3.58 (2H, s), 6.55-6.65
 5 (1H, m), 7.30-7.45 (2H, m), 7.60-7.70 (1H, m), 8.02 (1H, brs),
 8.05-8.15 (1H, m), 8.27 (1H, brs), 11.03 (1H, brs)

Compound 2-61

¹H-NMR (DMSO-d₆) δ ppm: 2.23 (6H, s), 3.57 (2H, s), 6.55-6.60
 (1H, m), 7.25-7.35 (1H, m), 7.40-7.50 (1H, m), 7.70-7.80 (1H,
 10 m), 8.01 (1H, brs), 8.25 (1H, brs), 8.36 (1H, d, J=9.1Hz), 11.05
 (1H, s)

MS (ESI, m/z): 362 (M+H)+

Compound 2-62

¹H-NMR (CDCl₃) δ ppm: 2.36 (6H, s), 3.62 (2H, s), 6.40-6.45 (1H,
 15 m), 6.50 (1H, brs), 6.95 (1H, brs), 6.95-7.05 (1H, m), 7.20-7.25
 (1H, m), 8.05-8.15 (1H, m), 10.80 (1H, brs)

Compound 2-63

¹H-NMR (CDCl₃) δ ppm: 0.85-0.95 (6H, m), 1.70-1.90 (1H, m),
 2.15-2.25 (2H, m), 2.32 (3H, s), 3.68 (2H, s), 5.99 (1H, brs),
 20 6.20-6.60 (2H, m), 7.15-7.40 (3H, m), 8.30-8.40 (1H, m), 10.76
 (1H, brs)

Compound 2-64

¹H-NMR (CDCl₃) δ ppm: 1.10 (6H, d, J=6.5Hz), 2.33 (3H, s), 2.90
 (1H, heptet, J=6.5Hz), 3.72 (2H, s), 5.99 (1H, brs), 6.30-6.60
 25 (2H, m), 7.10-7.40 (3H, m), 8.25-8.40 (1H, m), 10.74 (1H, brs)

Compound 2-65

¹H-NMR (CDCl₃) δ ppm: 1.18 (6H, t, J=7.2Hz), 2.67 (4H, q, J=7.2Hz),

3.90 (2H, s), 6.40-6.50 (1H, m), 6.75 (1H, brs), 6.91 (1H, brs),
7.10-7.40 (3H, m), 8.25-8.40 (1H, m), 10.84 (1H, brs)

Compound 2-66

¹H-NMR (DMSO-d₆) δ ppm: 0.90-1.10 (6H, m), 3.73 (2H, s), 6.57
5 (1H, brs), 7.25-7.35 (1H, m), 7.35-7.45 (1H, m), 7.70-7.80 (1H,
m), 8.01 (1H, brs), 8.25 (1H, brs), 8.30-8.45 (1H, m), 11.04
(1H, s)

MS (ESI, m/z): 390 (M+H)+

Compound 2-67

10 ¹H-NMR (DMSO-d₆) δ ppm: 1.04 (6H, t, J=6.9Hz), 2.40-2.60 (4H,
m), 3.73 (2H, s), 6.57 (1H, d, J=3.5Hz), 7.32 (1H, d, J=3.5Hz),
7.50-7.65 (1H, m), 7.90-8.00 (1H, m), 8.09 (1H, brs), 8.35 (1H,
brs), 10.98 (1H, s)

Compound 2-68

15 ¹H-NMR (CDCl₃) δ ppm: 1.06 (12H, d, J=6.5Hz), 3.10 (2H, heptet,
J=6.5Hz), 3.75 (2H, s), 5.78 (1H, brs), 6.30-6.50 (2H, m),
7.15-7.40 (3H, m), 8.30-8.45 (1H, m), 10.70 (1H, brs)

Compound 2-69

¹H-NMR (DMSO-d₆) δ ppm: 0.90-1.15 (12H, m), 2.90-3.15 (2H, m),
20 3.72 (2H, s), 6.45-6.60 (1H, m), 7.20-7.35 (1H, m), 7.35-7.50
(1H, m), 7.70-7.80 (1H, m), 7.99 (1H, brs), 8.24 (1H, brs),
8.30-8.45 (1H, m). 10.98 (1H, s)

MS (ESI, m/z): 418 (M+H)+

Compound 2-70

25 ¹H-NMR (CDCl₃+MeOD-d₄) δ ppm: 1.00-2.00 (10H, m), 2.35-2.45 (1H,
m), 2.37 (3H, s), 3.78 (2H, s), 6.40-6.50 (1H, m), 6.95-7.10
(1H, m), 7.20-7.35 (1H, m), 8.05-8.15 (1H, m)

Compound 2-71

MS (ESI, m/z): 460 (M+H)+

compound 2-72

MS (ESI, m/z): 394 (M+H)+

5 Compound 2-73

$^1\text{H-NMR}$ (CDCl_3) δ ppm: 1.14 (9H, s), 2.92 (2H, t, $J=6.0\text{Hz}$), 3.57 (2H, t, $J=6.0\text{Hz}$), 3.86 (2H, s), 6.30-6.40 (1H, m), 6.69 (1H, brs), 6.95-7.10 (1H, m), 7.20-7.30 (1H, m), 8.15-8.25 (1H, m), 10.88 (1H, brs)

10 Compound 2-74

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.00-1.65 (6H, m), 1.70-2.20 (4H, m), 3.15-3.90 (5H, m), 4.61 (2H, d, $J=3.8\text{Hz}$), 5.39 (1H, brs), 7.05 (1H, d, $J=3.5\text{Hz}$), 7.47 (1H, d, $J=3.5\text{Hz}$), 7.55-7.70 (1H, m), 7.85-7.95 (1H, m), 8.13 (1H, brs), 8.44 (1H, brs), 10.15 (1H, brs), 11.11 (1H, s)

Compound 2-75

MS (ESI, m/z): 408 (M+H)+

Compound 2-76

20 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.04 (3H, t, $J=7.1\text{Hz}$), 2.16 (6H, s), 2.30-2.45 (2H, m), 2.45-2.60 (4H, m), 3.77 (2H, s), 6.59 (1H, d, $J=3.5\text{Hz}$), 7.34 (1H, d, $J=3.5\text{Hz}$), 7.55-7.70 (1H, m), 7.90-8.00 (1H, m), 8.11 (1H, brs), 8.39 (1H, brs), 11.00 (1H, brs)

MS (ESI, m/z): 435 (M+H)+

Compound 2-77

25 $^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 2.30 (3H, s), 2.65-3.00 (4H, m), 3.71 (2H, s), 6.50-6.60 (1H, m), 7.10-7.40 (4H, m), 7.50-7.75 (2H, m), 7.85-8.00 (1H, m), 8.08 (1H, brs), 8.35-8.55 (2H, m)

Compound 2-78

¹H-NMR (DMSO-d₆) δ ppm: 2.55-2.65 (4H, m), 3.45-3.55 (4H, m),
3.83 (2H, s), 4.35-4.45 (2H, m), 6.55-6.65 (1H, m), 7.30-7.40
(1H, m), 7.55-7.65 (1H, m), 7.90-8.00 (1H, m), 8.14 (1H, brs),
5 8.41 (1H, brs), 10.99 (1H, brs)

Compound 2-79

¹H-NMR (DMSO-d₆) δ ppm: 2.70 (4H, t, J=6.0Hz), 3.24 (6H, s),
3.44 (4H, t, J=6.0Hz), 3.83 (2H, s), 6.59 (1H, d, J=3.5Hz), 7.33
(1H, d, J=3.5Hz), 7.55-7.70 (1H, m), 7.90-8.00 (1H, m), 8.10
10 (1H, brs), 8.38 (1H, brs), 10.99 (1H, s)

MS (ESI, m/z): 452 (M+H)+

Compound 2-80

¹H-NMR (DMSO-d₆) δ ppm: 1.04 (6H, d, J=6.4Hz), 1.70-1.85 (2H,
m), 2.70-2.80 (2H, m), 3.50-3.70 (4H, m), 6.60 (1H, d, J=3.3Hz),
15 7.32 (1H, d, J=3.3Hz), 7.35-7.50 (1H, m), 7.60-7.70 (1H, m),
7.95-8.10 (2H, m), 8.30 (1H, brs), 10.99 (1H, s)

MS (ESI, m/z): 416 (M+H)+

Compound 2-81

¹H-NMR (DMSO-d₆) δ ppm: 1.88 (1H, t, J=10.5Hz), 2.10-2.25 (1H,
20 m), 2.65-2.75 (1H, m), 2.82 (1H, d, J=10.9Hz), 3.25-3.60 (4H,
m), 3.64 (2H, s), 3.70-3.85 (1H, m), 4.70 (1H, t, J=5.7Hz), 6.61
(1H, d, J=3.5Hz), 7.34 (1H, d, J=3.5Hz), 7.50-7.65 (1H, m),
7.85-7.95 (1H, m), 8.10 (1H, brs), 8.40 (1H, brs), 10.95 (1H,
s)

25 MS (ESI, m/z): 436 (M+H)+

Compound 2-82

¹H-NMR (DMSO-d₆) δ ppm: 2.40-2.50 (4H, m), 3.50-3.65 (6H, m),

6.61 (1H, d, J=3.5Hz), 7.34 (1H, d, J=3.5Hz), 7.55-7.65 (1H, m), 7.85-7.95 (1H, m), 8.11 (1H, brs), 8.38 (1H, brs), 10.97 (1H, s)

MS (ESI, m/z): 406 (M+H)+

5 Compound 2-83

¹H-NMR (DMSO-d₆) δ ppm: 2.40-2.55 (4H, m), 3.55-3.70 (6H, m), 6.61 (1H, d, J=3.5Hz), 7.32 (1H, d, J=3.5Hz), 7.35-7.45 (1H, m), 7.60-7.70 (1H, m), 8.00 (1H, brs), 8.00-8.10 (1H, m), 8.27 (1H, brs), 11.00 (1H, s)

10 MS (ESI, m/z): 388 (M+H)+

Compound 2-84

¹H-NMR (DMSO-d₆) δ ppm: 2.40-2.55 (4H, m), 3.50-3.70 (6H, m), 6.55-6.65 (1H, m), 7.25-7.35 (1H, m), 7.40-7.50 (1H, m), 7.70-7.80 (1H, m), 8.00 (1H, brs), 8.25 (1H, brs), 8.34 (1H, d, J=9.1Hz), 11.02 (1H, s)

MS (ESI, m/z): 404 (M+H)+

Compound 2-85

¹H-NMR (DMSO-d₆) δ ppm: 1.41 (3H, d, J=6.3Hz), 1.55-1.70 (1H, m), 1.80-2.05 (2H, m), 2.15-2.30 (1H, m), 3.20-3.70 (3H, m), 4.40-4.55 (1H, m), 4.60-4.80 (1H, m), 7.01 (1H, d, J=3.5Hz), 7.46 (1H, d, J=3.5Hz), 7.55-7.70 (1H, m), 7.80-7.95 (1H, m), 8.13 (1H, brs), 8.44 (1H, brs), 10.62 (1H, brs), 11.09 (1H, s)

Compound 2-86

¹H-NMR (CDCl₃+MeOD-d₄) δ ppm: 1.15-1.65 (8H, m), 2.00-2.25 (2H, m), 2.55-3.05 (4H, m), 3.84 (2H, s), 6.35-6.45 (1H, m), 6.95-7.05 (1H, m), 7.15-7.30 (1H, m), 8.05-8.15 (1H, m)

Compound 2-87

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.40-1.55 (1H, m), 1.55-1.75 (2H, m),
 1.75-1.95 (1H, m), 2.30-2.60 (1H, m), 2.65-2.80 (1H, m),
 2.85-3.05 (1H, m), 3.15-3.45 (5H, m), 3.55-3.75 (1H, m),
 4.00-4.15 (1H, m), 6.50-6.60 (1H, m), 7.25-7.35 (1H, m),
 5 7.55-7.65 (1H, m), 7.85-8.00 (1H, m), 8.12 (1H, brs), 8.38 (1H,
 brs), 10.97 (1H, s)

MS (ESI, m/z): 434 (M+H)+

Compound 2-88

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.60-2.30 (4H, m), 3.20-3.90 (8H, m),
 10 4.45-4.75 (2H, m), 7.01 (1H, d, J=3.5Hz), 7.46 (1H, d, J=3.5Hz),
 7.55-7.70 (1H, m), 7.80-7.90 (1H, m), 8.12 (1H, brs), 8.44 (1H,
 brs), 10.76 (1H, brs), 11.09 (1H, s)

Compound 2-89

MS (ESI, m/z): 420 (M+H)+

15 Compound 2-90

MS (ESI, m/z): 420 (M+H)+

Compound 2-91

$^1\text{H-NMR}$ (DMSO-d_6) δ ppm: 1.90-2.05 (2H, m), 2.30 (2H, t, J=8.1Hz),
 3.45 (2H, t, J=7.1Hz), 4.51 (2H, s), 6.55-6.65 (1H, m), 7.30-7.40
 20 (1H, m), 7.55-7.70 (1H, m), 7.85-8.00 (1H, m), 8.12 (1H, brs),
 8.39 (1H, brs), 10.99 (1H, s)

MS (ESI, m/z): 404 (M+H)+

Compound 2-92

MS (ESI, m/z): 433 (M+H)+

25 Compound 2-93

MS (ESI, m/z): 433 (M+H)+

Compound 2-94

MS (ESI, m/z): 467 (M+H)+

Compound 2-95

MS (ESI, m/z): 467 (M+H)+

Compound 2-96

- 5 $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 1.30-1.45 (2H, m), 1.45-1.60 (4H, m),
2.35-2.50 (4H, m), 3.58 (2H, s), 6.56 (1H, d, $J=3.5\text{Hz}$), 7.30
(1H, d, $J=3.5\text{Hz}$), 7.35-7.45 (1H, m), 7.60-7.70 (1H, m), 8.00
(1H, brs), 8.00-8.15 (1H, m), 8.27 (1H, brs), 10.98 (1H, brs)

MS (ESI, m/z): 386 (M+H)+

- 10 Compound 2-97

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 1.30-1.45 (2H, m), 1.45-1.60 (4H, m),
2.30-2.50 (4H, m), 3.59 (2H, s), 6.55-6.60 (1H, m), 7.30-7.40
(1H, m), 7.55-7.65 (1H, m), 7.85-7.95 (1H, m), 8.10 (1H, brs),
8.37 (1H, brs), 10.96 (1H, s)

- 15 MS (ESI, m/z): 404 (M+H)+

Compound 2-98

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 1.05-1.65 (9H, m), 2.10-2.35 (2H, m),
2.75-2.90 (1H, m), 3.81 (2H, brs), 6.55-6.65 (1H, m), 7.30-7.50
(2H, m), 7.60-7.75 (1H, m), 8.00-8.20 (2H, m), 8.31 (1H, brs),
20 11.05 (1H, s)

MS (ESI, m/z): 400 (M+H)+

Compound 2-99

- $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 1.05-1.30 (5H, m), 1.30-1.65 (4H, m),
2.10-2.30 (2H, m), 2.75-2.85 (1H, m), 3.78 (2H, s), 6.58 (1H,
25 d, $J=3.4\text{Hz}$), 7.33 (1H, d, $J=3.4\text{Hz}$), 7.55-7.70 (1H, m), 7.90-8.00
(1H, m), 8.12 (1H, brs), 8.38 (1H, brs), 11.00 (1H, s)

MS (ESI, m/z): 418 (M+H)+

Compound 2-100

¹H-NMR (DMSO-d₆) δ ppm: 0.85-1.00 (3H, m), 1.30-2.05 (7H, m),
2.10-2.25 (1H, m), 2.90-3.60 (3H, m), 4.40-4.70 (2H, m),
6.95-7.10 (1H, m), 7.40-7.55 (1H, m), 7.55-7.70 (1H, m),
5 7.80-7.95 (1H, m), 8.05-8.20 (1H, m), 8.35-8.50 (1H, m), 10.43
(1H, brs), 11.00-11.20 (1H, m)

Compound 2-101

¹H-NMR (DMSO-d₆) δ ppm: 1.10-1.30 (9H, m), 1.45-1.65 (3H, m),
2.25-2.40 (2H, m), 3.97 (2H, s), 6.50-6.65 (1H, m), 7.32 (1H,
10 d, J=3.4Hz), 7.35-7.45 (1H, m), 7.60-7.70 (1H, m), 8.02 (1H,
brs), 8.10-8.20 (1H, m), 8.26 (1H, brs), 11.02 (1H, s)

MS (ESI, m/z): 414 (M+H)+

Compound 2-102

¹H-NMR (DMSO-d₆) δ ppm: 1.00-1.35 (9H, m), 1.45-1.65 (3H, m),
15 2.25-2.50 (2H, m), 3.98 (2H, brs), 6.50-6.65 (1H, m), 7.25-7.40
(1H, m), 7.40-7.50 (1H, m), 7.70-7.80 (1H, m), 8.01 (1H, brs),
8.25 (1H, brs), 8.35-8.50 (1H, m), 11.05 (1H, s)

MS (ESI, m/z): 430 (M+H)+

Compound 2-103

20 ¹H-NMR (DMSO-d₆) δ ppm: 1.10-1.30 (9H, m), 1.45-1.65 (3H, m),
2.25-2.40 (2H, m), 3.97 (2H, s), 6.59 (1H, d, J=3.6Hz), 7.34
(1H, d, J=3.6Hz), 7.55-7.70 (1H, m), 7.90-8.05 (1H, m), 8.12
(1H, brs), 8.38 (1H, brs), 11.00 (1H, s)

MS (ESI, m/z): 432 (M+H)+

25 Compound 2-104

¹H-NMR (DMSO-d₆) δ ppm: 0.40-0.55 (1H, m), 0.81 (6H, d, J=6.3Hz),
1.45-1.70 (5H, m), 2.75-2.90 (2H, m), 3.59 (2H, s), 6.50-6.60

(1H, m), 7.25-7.35 (1H, m), 7.35-7.45 (1H, m), 7.60-7.70 (1H, m), 7.95-8.10 (2H, m), 8.29 (1H, brs), 10.97 (1H, s)

MS (ESI, m/z): 414 (M+H)+

Compound 2-105

5 MS (ESI, m/z): 452 (M+H)+

Compound 2-106

¹H-NMR (DMSO-d₆) δ ppm: 1.30-1.50 (2H, m), 1.65-1.80 (2H, m), 2.05-2.25 (2H, m), 2.65-2.85 (2H, m), 3.35-3.50 (1H, m), 3.60 (2H, s), 4.57 (1H, d, J=4.0Hz), 6.58 (1H, d, J=3.5Hz), 7.33 (1H, d, J=3.5Hz), 7.55-7.65 (1H, m), 7.85-7.95 (1H, m), 8.12 (1H, brs), 8.38 (1H, brs), 10.96 (1H, s)

MS (ESI, m/z): 420 (M+H)+

Compound 2-107

MS (ESI, m/z): 434 (M+H)+

15 Compound 2-108

¹H-NMR (DMSO-d₆) δ ppm: 0.75-0.95 (1H, m), 1.35-1.80 (5H, m), 1.95-2.05 (1H, m), 2.75-2.85 (1H, m), 2.85-2.95 (1H, m), 3.15-3.40 (2H, m), 3.59 (2H, s), 4.41 (1H, t, J=5.3Hz), 6.55-6.60 (1H, m), 7.30-7.35 (1H, m), 7.35-7.45 (1H, m), 7.60-7.70 (1H, m), 7.95-8.10 (2H, m), 8.30 (1H, brs), 10.98 (1H, s)

MS (ESI, m/z): 416 (M+H)+

Compound 2-109

¹H-NMR (CDCl₃+MeOD-d₄) δ ppm: 1.25-1.85 (5H, m), 2.10-2.25 (2H, m), 2.95-3.05 (2H, m), 3.30-3.50 (2H, m), 3.69 (2H, s), 6.45 (1H, d, J=3.5Hz), 7.00-7.10 (1H, m), 7.26 (1H, d, J=3.5Hz), 8.05-8.15 (1H, m)

Compound 2-110

- ¹H-NMR (DMSO-d₆) δ ppm: 0.75-0.95 (1H, m), 1.35-1.80 (5H, m), 1.90-2.05 (1H, m), 2.70-2.85 (1H, m), 2.85-2.95 (1H, m), 3.10-3.40 (2H, m), 3.59 (2H, s), 4.41 (1H, t, J=5.2Hz), 6.50-6.65 (1H, m), 7.25-7.40 (1H, m), 7.50-7.70 (1H, m), 7.85-7.95 (1H, m), 8.10 (1H, brs), 8.38 (1H, brs), 10.94 (1H, s)
- 5 MS (ESI, m/z): 434 (M+H)+
Compound 2-111
- MS (ESI, m/z): 448 (M+H)+
Compound 2-112
- 10 MS (ESI, m/z): 447 (M+H)+
Compound 2-113
- ¹H-NMR (DMSO-d₆) δ ppm: 1.20-1.55 (2H, m), 1.55-1.75 (2H, m), 1.95-2.15 (2H, m), 2.25-2.40 (1H, m), 2.75-2.90 (2H, m), 3.61 (2H, s), 6.59 (1H, d, J=3.2Hz), 6.78 (1H, brs), 7.25-7.40 (2H, m), 7.55-7.65 (1H, m), 7.85-7.95 (1H, m), 8.10 (1H, brs), 8.40 (1H, brs), 10.94 (1H, brs)
- 15 MS (ESI, m/z): 447 (M+H)+
Compound 2-114
- ¹H-NMR (DMSO-d₆) δ ppm: 1.55-1.70 (4H, m), 3.65 (2H, s), 3.84 (4H, s), 6.55-6.65 (1H, m), 7.30-7.40 (1H, m), 7.55-7.65 (1H, m), 7.85-7.95 (1H, m), 8.12 (1H, brs), 8.38 (1H, brs), 10.97 (1H, brs)
- 20 MS (ESI, m/z): 462 (M+H)+
Compound 2-115
- ¹H-NMR (DMSO-d₆) δ ppm: 1.55-1.70 (4H, m), 2.45-2.60 (4H, m), 3.65 (2H, s), 3.84 (4H, s), 6.55-6.65 (1H, m), 7.20-7.40 (1H, m), 7.42 (1H, dd, J=1.9, 8.8Hz), 7.70-7.80 (1H, m), 8.01 (1H, m), 8.12 (1H, brs), 8.38 (1H, brs), 10.97 (1H, brs)
- 25

brs), 8.25 (1H, brs), 8.34 (1H, d, J=8.8Hz), 11.02 (1H, s)

MS (ESI, m/z): 460 (M+H)+

Compound 2-116

¹H-NMR (DMSO-d₆) δ ppm: 1.45-2.00 (8H, m), 3.10-3.60 (4H, m),
 5 4.54 (2H, d, J=4.1Hz), 7.02 (1H, d, J=3.8Hz), 7.46 (1H, d,
 J=3.8Hz), 7.55-7.70 (1H, m), 7.80-7.90 (1H, m), 8.12 (1H, brs),
 8.43 (1H, brs), 10.83 (1H, brs), 11.10 (1H, s)

Compound 2-117

¹H-NMR (DMSO-d₆) δ ppm: 1.40-1.65 (10H, m), 2.55-2.70 (4H, m),
 10 3.75 (2H, s), 6.57 (1H, d, J=3.4Hz), 7.33 (1H, d, J=3.4Hz),
 7.55-7.70 (1H, m), 7.90-8.00 (1H, m), 8.11 (1H, brs), 8.38 (1H,
 brs), 10.99 (1H, s)

MS (ESI, m/z): 432 (M+H)+

Compound 2-118

15 ¹H-NMR (DMSO-d₆) δ ppm: 2.10-2.70 (11H, m), 3.61 (2H, s), 6.55-6.65
 (1H, m), 7.25-7.40 (1H, m), 7.55-7.65 (1H, m), 7.85-7.95 (1H,
 m), 8.10 (1H, brs), 8.38 (1H, brs), 10.96 (1H, brs)

MS (ESI, m/z): 419 (M+H)+

Compound 2-119

20 ¹H-NMR (DMSO-d₆) δ ppm: 2.16 (3H, s), 2.20-2.60 (8H, m), 3.62
 (2H, s), 6.58 (1H, d, J=3.5Hz), 7.31 (1H, d, J=3.5Hz), 7.35-7.45
 (1H, m), 7.60-7.70 (1H, m), 7.95-8.15 (2H, m), 8.27 (1H, brs),
 10.99 (1H, s)

MS (ESI, m/z): 401 (M+H)+

25 Compound 2-120

¹H-NMR (DMSO-d₆) δ ppm: 0.85-1.05 (6H, m), 2.20-2.70 (9H, m),
 3.61 (2H, s), 6.58 (1H, d, J=3.5Hz), 7.33 (1H, d, J=3.5Hz),

7.55-7.65 (1H, m), 7.85-7.95 (1H, m), 8.08 (1H, brs), 8.35 (1H, brs), 10.95 (1H, s)

MS (ESI, m/z): 447 (M+H)+

Compound 2-121

5 MS (ESI, m/z): 501 (M+H)+

Compound 2-122

MS (ESI, m/z): 477 (M+H)+

Compound 2-123

MS (ESI, m/z): 490 (M+H)+

10 Compound 2-124

MS (ESI, m/z): 532 (M+H)+

Compound 2-125

¹H-NMR (DMSO-d₆) δ ppm: 1.31 (3H, t, J=6.9Hz), 2.63 (4H, brs), 3.01 (4H, brs), 3.70 (2H, s), 3.99 (2H, q, J=6.9Hz), 6.63 (1H, d, J=3.1Hz), 6.80-6.95 (4H, m), 7.33 (1H, d, J=3.1Hz), 7.35-7.50 (1H, m), 7.70-7.80 (1H, m), 7.99 (1H, brs), 8.25 (1H, brs), 8.34 (1H, d, J=9.0Hz), 11.04 (1H, s)

MS (ESI, m/z): 523 (M+H)+

Compound 2-126

20 ¹H-NMR (DMSO-d₆) δ ppm: 1.32 (3H, t, J=7.0Hz), 2.55-2.70 (4H, m), 2.90-3.10 (4H, m), 3.70 (2H, s), 3.99 (2H, q, J=7.0Hz), 6.60-6.70 (1H, m), 6.80-6.95 (4H, m), 7.30-7.45 (2H, m), 7.60-7.70 (1H, m), 8.00 (1H, brs), 8.05-8.15 (1H, m), 8.27 (1H, brs), 11.02 (1H, brs)

25 Compound 2-127

¹H-NMR (DMSO-d₆) δ ppm: 2.20-2.60 (8H, m), 3.45 (2H, s), 3.62 (2H, s), 6.50-6.60 (1H, m), 7.15-7.35 (6H, m), 7.35-7.45 (1H,

m), 7.60–7.70 (1H, m), 7.90–8.15 (2H, m), 8.26 (1H, brs), 10.98 (1H, s)

MS (ESI, m/z): 477 (M+H)+

Compound 2-128

5 $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 2.20–2.60 (4H, m), 3.45 (2H, s), 3.62 (2H, s), 6.50–6.60 (1H, m), 7.20–7.40 (6H, m), 7.40–7.50 (1H, m), 7.70–7.80 (1H, m), 7.99 (1H, brs), 8.24 (1H, brs), 8.30–8.40 (1H, m), 11.01 (1H, s)

MS (ESI, m/z): 493 (M+H)+

10 Compound 2-129

MS (ESI, m/z): 495 (M+H)+

Compound 2-130

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 2.30–2.60 (8H, m), 3.51 (2H, s), 3.62 (2H, s), 6.57 (1H, d, $J=3.1\text{Hz}$), 7.10–7.20 (2H, m), 7.25–7.35 (2H, m), 7.35–7.45 (1H, m), 7.55–7.65 (1H, m), 7.85–7.95 (1H, m), 8.08 (1H, brs), 8.34 (1H, brs), 10.95 (1H, s)

15

MS (ESI, m/z): 513 (M+H)+

Compound 2-131

$^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 2.25–2.60 (8H, m), 3.48 (2H, s), 3.63 (2H, s), 6.58 (1H, d, $J=3.7\text{Hz}$), 7.00–7.20 (3H, m), 7.30–7.40 (2H, m), 7.50–7.65 (1H, m), 7.85–7.95 (1H, m), 8.07 (1H, brs), 8.34 (1H, brs), 10.95 (1H, s)

20

MS (ESI, m/z): 513 (M+H)+

Compound 2-132

25 $^1\text{H-NMR}$ (DMSO- d_6) δ ppm: 2.20–2.60 (8H, m), 3.43 (2H, s), 3.62 (2H, s), 6.50–6.65 (1H, m), 7.05–7.20 (2H, m), 7.20–7.40 (3H, m), 7.55–7.65 (1H, m), 7.85–7.95 (1H, m), 8.07 (1H, brs), 8.34

(1H, brs), 10.94 (1H, s)

MS (ESI, m/z): 513 (M+H)+

Compound 2-133

¹H-NMR (DMSO-d₆) δ ppm: 2.25-2.60 (8H, m), 3.42 (2H, brs), 3.62
5 (2H, s), 3.73 (3H, s), 6.58 (1H, d, J=3.5Hz), 6.75-6.95 (3H, m), 7.15-7.30 (1H, m), 7.32 (1H, d, J=3.5Hz), 7.55-7.65 (1H, m), 7.85-7.95 (1H, m), 8.07 (1H, brs), 8.34 (1H, brs), 10.95 (1H, s)

MS (ESI, m/z): 525 (M+H)+

10 Compound 2-134

¹H-NMR (DMSO-d₆) δ ppm: 2.20-2.65 (8H, m), 3.25-3.45 (2H, m), 3.62 (2H, s), 3.72 (3H, s), 6.57 (1H, brs), 6.80-6.95 (2H, m), 7.10-7.25 (2H, m), 7.32 (1H, brs), 7.50-7.65 (1H, m), 7.85-8.00 (1H, m), 8.07 (1H, brs), 8.34 (1H, brs), 10.94 (1H, s)

15 MS (ESI, m/z): 525 (M+H)+

Compound 2-135

¹H-NMR (DMSO-d₆) δ ppm: 2.25-2.60 (8H, m), 3.38 (2H, s), 3.62 (2H, s), 3.71 (6H, s), 6.30-6.40 (1H, m), 6.40-6.55 (2H, m), 6.58 (1H, d, J=3.4Hz), 7.32 (1H, d, J=3.4Hz), 7.55-7.65 (1H, m),
20 m), 7.85-7.95 (1H, m), 8.08 (1H, brs), 8.35 (1H, brs), 10.95 (1H, s)

MS (ESI, m/z): 555 (M+H)+

Compound 2-136

¹H-NMR (DMSO-d₆) δ ppm: 2.25-2.65 (8H, m), 3.49 (2H, s), 3.63
25 (2H, s), 6.50-6.65 (1H, m), 7.30-7.45 (2H, m), 7.50-7.80 (2H, m), 7.85-8.00 (1H, m), 8.00-8.20 (1H, m), 8.25-8.55 (3H, m), 10.95 (1H, s)

MS (ESI, m/z): 496 (M+H)+

Compound 2-137

¹H-NMR (DMSO-d₆) δ ppm: 2.30-2.60 (8H, m), 3.50 (2H, s), 3.64
(2H, s), 6.55-6.65 (1H, m), 7.25-7.40 (3H, m), 7.55-7.65 (1H,
5 m), 7.85-8.00 (1H, m), 8.00-8.15 (1H, m), 8.25-8.45 (1H, m),
8.45-8.60 (2H, m), 10.95 (1H, s)

MS (ESI, m/z): 496 (M+H)+

Compound 2-138

MS (ESI, m/z): 539 (M+H)+

10 Compound 2-139

MS (ESI, m/z): 571 (M+H)+

Compound 2-140

¹H-NMR (DMSO-d₆) δ ppm: 1.98 (3H, s), 2.35-2.55 (4H, m), 3.35-3.55
(4H, m), 3.68 (2H, s), 6.61 (1H, d, J=3.5Hz), 7.34 (1H, d, J=3.5Hz),
15 7.55-7.70 (1H, m), 7.85-7.95 (1H, m), 8.11 (1H, brs), 8.38 (1H,
brs), 10.97 (1H, s)

Compound 2-141

¹H-NMR (DMSO-d₆) δ ppm: 2.35-2.65 (4H, m), 3.50-3.80 (4H, m),
6.55-6.70 (1H, m), 7.30-7.50 (6H, m), 7.55-7.70 (1H, m),
20 7.85-8.00 (1H, m), 8.12 (1H, brs), 8.39 (1H, brs), 10.97 (1H,
s)

MS (ESI, m/z): 509 (M+H)+

Compound 2-142

¹H-NMR (DMSO-d₆) δ ppm: 1.38 (9H, s), 2.35-2.50 (4H, m), 3.66
25 (2H, s), 6.61 (1H, d, J=3.4Hz), 7.34 (1H, d, J=3.4Hz), 7.55-7.70
(1H, m), 7.85-8.00 (1H, m), 8.11 (1H, brs), 8.38 (1H, brs), 10.97
(1H, s)

MS (ESI, m/z): 505 (M+H)+

Compound 2-143

¹H-NMR (CDCl₃) δ ppm: 2.35-2.80 (12H, m), 3.67 (2H, s), 5.12
(2H, s), 5.70 (1H, brs), 6.35-6.45 (1H, m), 6.48 (1H, brs),
5 6.95-7.10 (1H, m), 7.20-7.45 (6H, m), 8.10-8.25 (1H, m), 10.72
(1H, brs)

Compound 2-144

¹H-NMR (DMSO-d₆) δ ppm: 3.45-3.60 (2H, m), 3.60-3.80 (2H, m),
4.00-4.20 (2H, m), 4.67 (2H, s), 5.11 (2H, m), 6.62 (1H, d,
10 J=3.1Hz), 7.25-7.45 (6H, m), 7.50-7.65 (1H, m), 7.85-8.00 (1H,
m), 8.08 (1H, brs), 8.36 (1H, brs), 11.01 (1H, s)

MS (ESI, m/z): 553 (M+H)+

Compound 2-145

¹H-NMR (DMSO-d₆) δ ppm: 1.60-1.80 (3H, m), 2.20-2.80 (10H, m),
15 3.75 (2H, s), 6.57 (1H, d, J=3.5Hz), 7.32 (1H, d, J=3.5Hz),
7.55-7.65 (1H, m), 7.85-8.00 (1H, m), 8.10 (1H, brs), 8.40 (1H,
brs), 11.00 (1H, brs)

Example 4

5-Fluoro-3-(4-hydroxy-3-morpholin-4-ylmethylbenzoylamino)-benzofuran-2-carboxamide (compound 3-1)

3-(4-Benzyloxy-3-morpholin-4-ylmethylbenzoylamino)-5-fluorobenzofuran-2-carboxamide (compound 1-202, 0.026g) was dissolved in tetrahydrofuran (3mL) and 10% palladium on carbon (50% wet, 0.025g) was added to the solution, and the mixture was stirred at room temperature under hydrogen atmosphere for 6 hours. After the catalyst was removed by filtration, the filtrate was concentrated under reduced pressure. The residue was purified by a SCX ion-exchanged column conditioned with methanol (argonaut 1g, eluent: methanol) to give the title compound (0.01g).

¹H-NMR (CDCl₃) δ ppm: 2.45-2.80 (4H, m), 2.96 (1H, brs), 3.65-4.00 (6H, m), 5.78 (1H, brs), 6.43 (1H, brs), 6.90-7.00 (1H, m), 7.15-7.25 (1H, m), 7.35-7.45 (1H, m), 7.70-7.80 (1H, m), 7.85-7.95 (1H, m), 8.30-8.40 (1H, m), 10.73 (1H, brs)

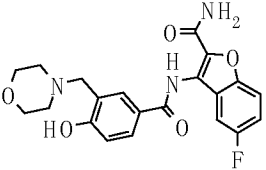
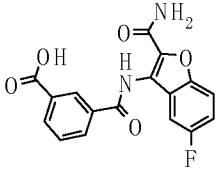
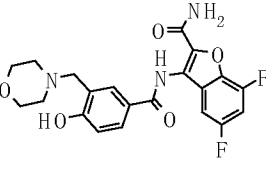
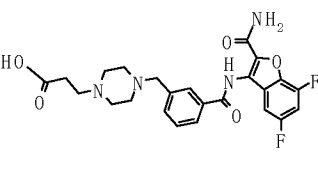
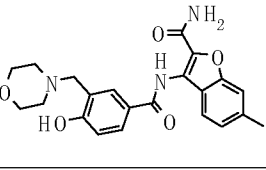
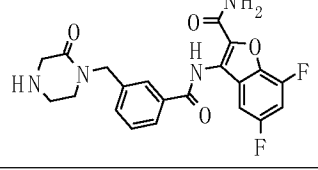
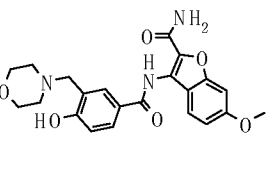
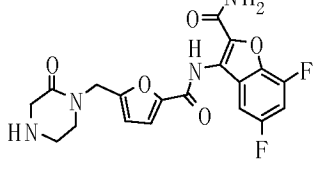
Compounds 3-2 to 3-8 were prepared in a manner similar to those as described in Example 4 using

3-(4-benzyloxy-3-morpholin-4-ylmethylbenzoylamino)-5,7-difluorobenzofuran-2-carboxamide (compound 1-203),
 3-(4-benzyloxy-3-morpholin-4-ylmethylbenzoylamino)-6-methylbenzofuran-2-carboxamide (compound 1-204),
 3-(4-benzyloxy-3-morpholin-4-ylmethylbenzoylamino)-6-methoxybenzofuran-2-carboxamide (compound 1-205), benzyl N-(2-carbamoyl-5-fluorobenzofuran-3-yl)isophthalamate

(compound 1-235), benzyl 3-{4-[3-(2-carbamoyl-5,7-difluorobenzofuran-3-ylcarbamoyl)benzyl]piperazin-1-yl}propionate (compound 2-53), benzyl 4-[3-(2-carbamoyl-5,7-difluorobenzofuran-3-ylcarbamoyl)benzyl]-3-oxopiperazine-1-carboxylate (compound 2-55) and benzyl 4-[5-(2-carbamoyl-5,7-difluorobenzofuran-3-ylcarbamoyl)furan-2-ylmethyl]-3-oxopiperazine-1-carboxylate (compound 2-144) instead of 3-(4-benzyloxy-3-morpholin-4-ylmethylbenzoylamino)-5-fluorobenzofuran-2-carboxamide (compound 1-202).

These were illustrated in table 4.

Table 4

Compound No.	Structure	Compound No.	Structure
3-1		3-5	
3-2		3-6	
3-3		3-7	
3-4		3-8	

The physical data of compounds 3-2 to 3-8 were described below.

Compound 3-2

¹H-NMR (CDCl₃+MeOD-d₄) δ ppm: 2.50-2.80 (4H, m), 3.70-3.85 (4H, m), 3.83 (2H, s), 6.90-7.10 (2H, m), 7.70-7.90 (2H, m), 8.05-8.15 (1H, m)

Compound 3-3

¹H-NMR (CDCl₃) δ ppm: 2.50 (3H, s), 2.40-2.80 (4H, m), 3.70-3.85 (4H, m), 3.82 (2H, s), 5.54 (1H, brs), 6.36 (1H, brs), 6.90-6.95 (1H, m), 7.10-7.25 (2H, m), 7.75-7.80 (1H, m), 7.85-7.95 (1H, m), 8.45-8.55 (1H, m), 10.80 (1H, brs)

Compound 3-4

¹H-NMR (CDCl₃) δ ppm: 2.40-2.80 (4H, m), 2.99 (1H, brs), 3.70-3.85 (6H, m), 3.89 (3H, s), 5.52 (1H, brs), 6.30 (1H, brs), 6.85-7.00 (3H, m), 7.70-7.80 (1H, m), 7.85-7.95 (1H, m), 8.50-8.60 (1H, m), 10.86 (1H, brs)

Compound 3-5

¹H-NMR (DMSO-d₆) δ ppm: 7.35-7.50 (1H, m), 7.62-7.80 (2H, m), 7.90-8.09 (2H, m), 8.15-8.35 (3H, m), 8.60 (1H, s), 11.13 (1H, s), 12.80-13.78 (1H, br)

Compound 3-6

¹H-NMR (MeOD-d₄) δ ppm: 2.40-2.85 (6H, m), 2.95-3.20 (4H, m), 3.70 (2H, s), 7.15-7.30 (1H, m), 7.45-7.65 (2H, m), 7.85-8.10 (3H, m)

Compound 3-7

¹H-NMR (DMSO-d₆) δ ppm: 2.85-2.95 (2H, m), 3.15-3.30 (2H, m), 4.62 (2H, s), 7.50-7.70 (3H, m), 7.75-8.00 (3H, m), 8.10 (1H,

brs), 8.37 (1H, brs), 10.99 (1H, s)

MS (ESI, m/z): 429 (M+H)+

Compound 3-8

¹H-NMR (DMSO-d₆) δ ppm: 2.60-2.85 (1H, m), 2.93 (2H, t, J=5.5Hz),
 5 3.29 (2H, s), 3.39 (2H, t, J=5.5Hz), 4.62 (2H, s), 6.60 (1H,
 d, J=3.4Hz), 7.34 (1H, d, J=3.4Hz), 7.50-7.65 (1H, m), 7.85-8.00
 (1H, m), 8.11 (1H, brs), 8.39 (1H, brs), 10.99 (1H, s)

MS (ESI, m/z): 419 (M+H)+

10 Example 5

5,7-Difluoro-3-[5-(4-methylpiperazine-1-carbonyl)furan-2-carbonyl]aminobenzofuran-2-carboxamide (compound 4-1)

To a solution of furan-2,5-dicarbonyl dichloride (0.073g) in tetrahydrofuran (3mL) were added 3-amino-5,7-difluoro-
 15 benzofuran-2-carboxamide (0.08g) and N,N-dimethylaniline (0.096mL) successively under ice-cooling, and the resulting mixture was stirred at room temperature overnight. 1-Methylpiperazine (0.085mL) was added to the mixture, and the resulting mixture was stirred for 3 hours. Water and ethyl acetate were
 20 added to the reaction mixture. The organic layer was separated and washed with water. The organic layer was extracted with 2mol/L hydrochloric acid, and the aqueous layer was washed with ethyl acetate. Then, the aqueous layer was made alkaline with a saturated aqueous solution of sodium carbonate, and was extracted
 25 with ethyl acetate. The organic layer was separated, and washed with water, a saturated aqueous solution of sodium carbonate and brine, and dried over anhydrous magnesium sulfate. The

solvent was removed under reduced pressure, and the residue was purified by column chromatography on aminopropylated silica gel (eluent: hexane/ethyl acetate = 9/1 - 0/1) to give the title compound (0.004g). The structural formula was illustrated in table 5.

¹H-NMR (CDCl₃+MeOD-d₄) δ ppm: 2.30-2.65 (6H, m), 2.37 (3H, s), 3.95-4.20 (2H, m), 7.00-7.10 (1H, m), 7.20 (1H, d, J=3.5Hz), 7.35 (1H, d, J=3.5Hz), 8.10-8.25 (1H, m)

3-[5-(4-Benzylpiperazine-1-carbonyl)furan-2-carbonyl] amino-5,7-difluorobenzofuran-2-carboxamide (compounds 4-2) was prepared in a manner similar to those as described in Example 5 using 1-benzylpiperazine instead of 1-methylpiperazine. The structural formula was illustrated in table 5.

¹H-NMR (CDCl₃+DMSO-d₆) δ ppm: 2.50-2.70 (4H, m), 2.88 (2H, s), 3.55-3.65 (2H, m), 3.90-4.10 (2H, m), 6.04 (1H, brs), 6.69 (1H, brs), 7.00-7.40 (8H, m), 8.15-8.25 (1H, m)

Example 6

5-(2-Carbamoyl-5,7-difluorobenzofuran-3-ylcarbamoyl)furan-2-carboxylic acid (compound 5-1)

To a solution of furan-2,5-dicarbonyl dichloride (0.073g) in tetrahydrofuran (3mL) were added 3-amino-5,7-difluorobenzofuran-2-carboxamide (0.08g) and N,N-dimethylaniline (0.096mL) successively under ice-cooling, and the resulting mixture was stirred at room temperature overnight. A saturated aqueous solution of sodium carbonate and ethyl acetate were added

to the reaction mixture, and the aqueous layer was separated and washed with ethyl acetate. To the aqueous layer was added 2mol/L hydrochloric acid, and the resulting precipitate was collected by filtration. The precipitate was washed with water and methanol to give the title compound (0.018g). The structural formula was illustrated in table 5.

¹H-NMR (DMSO-d₆) δppm: 7.40 (1H, d, J=3.8Hz), 7.47 (1H, d, J=3.8Hz), 7.55-7.65 (1H, m), 7.75-7.90 (1H, m), 8.09 (1H, brs), 8.36 (1H, brs), 11.09 (1H, brs), 13.69 (1H, brs)

Example 7

5-Fluoro-3-[3-(1-hydroxyethyl)benzoylamino]benzofuran-2-carboxamide (compound 6-1)

Sodium borohydride (0.02g) was added to a mixture of 3-(3-acetylbenzoylamino)-5-fluorobenzofuran-2-carboxamide (compound 1-236, 0.045g), tetrahydrofuran (5mL) and methanol (3mL), and the resulting mixture was stirred at room temperature for 2 hours. Water was added to the reaction mixture and the mixture was stirred for another 1 hour. The mixture was concentrated under reduced pressure. To the residue were added water and ethyl acetate. The organic layer was separated, washed with brine, and dried over anhydrous magnesium sulfate. The solvent was removed under reduced pressure, and the residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate = 3/1 - 0/1) to give the title compound (0.038g). The structural formula was illustrated in the table 5.

¹H-NMR (CDCl₃) δ ppm: 1.54 (3H, d, J=6.5Hz), 3.47 (1H, d, J=4.0Hz), 4.95-5.05 (1H, m), 6.48 (1H, brs), 6.65 (1H, brs), 7.15-7.25 (1H, m), 7.35-7.55 (2H, m), 7.60-7.70 (1H, m), 7.85-7.95 (1H, m), 8.00-8.10 (1H, m), 8.30-8.40 (1H, m), 10.96 (1H, brs)

5

5-Fluoro-3-[5-(1-hydroxyethyl)furan-2-carbonyl]-aminobenzofuran-2-carboxamide (compound 6-2) was prepared in a manner similar to those as described in Example 7-1 using 3-(5-acetylfuran-2-carbonyl)amino-5-fluorobenzofuran-2-carboxamide (compound 1-304) instead of 3-(3-acetylbenzoylamino)-5-fluorobenzofuran-2-carboxamide (compound 1-236). The structural formula was illustrated in table 5.

¹H-NMR (CDCl₃) δ ppm: 1.63 (3H, d, J=6.6Hz), 3.34 (1H, d, J=2.5Hz), 4.90-5.05 (1H, m), 6.09 (1H, brs), 6.44 (1H, d, J=3.5Hz), 6.51 (1H, brs), 7.15-7.25 (1H, m), 7.23 (1H, d, J=3.5Hz), 7.30-7.40 (1H, m), 8.30-8.40 (1H, m), 10.88 (1H, brs)

Example 8

5,7-Difluoro-3-[3-(1H-tetrazol-5-yl)benzoylamino]benzofuran-2-carboxamide (compound 7-1)

A mixture of 3-(3-cyanobenzoylamino)-5,7-difluorobenzofuran-2-carboxamide (compound 1-206, 0.12g), sodium azide (0.093g) trimethylamine hydrochloride (0.101g) and 1-methyl-2-pyrrolidone (3mL) was stirred at 130°C overnight. Water (7mL) was added to the reaction mixture, and the mixture was acidified with the addition of concentrated hydrochloric acid. The mixture

was stirred at room temperature for 1 hour, and the resulting precipitate was collected by filtration. The crude product was purified by column chromatography on silica gel (eluent: dichloromethane/methanol=19/1 - 5/1) to give the title compound
 5 (0.046g). The structural formula was illustrated in the table 5.

¹H-NMR (DMSO-d₆) δ ppm: 7.55-7.65 (1H, m), 7.70-7.90 (2H, m), 8.08 (1H, brs), 8.10-8.20 (1H, m), 8.25-8.35 (1H, m), 8.34 (1H, brs), 8.65-8.75 (1H, m), 11.08 (1H, brs)

10

Example 9

7-Ethoxy-5-fluoro-3-(4-fluorobenzoylamino)benzofuran-2-carboxamide (compound 8-1)

To a mixture of 5-fluoro-3-(4-fluorobenzoylamino)-
 15 7-hydroxybenzofuran-2-carboxamide (compound 1-178, 0.08g), potassium carbonate (0.1g) and N,N-dimethylformamide (2mL) was added ethyl iodide (0.029mL) at room temperature, and the resulting mixture was stirred at 60°C overnight. Water was added to the reaction mixture, and the resulting precipitate was
 20 collected by filtration to give the title compound (0.04g). The structural formula was illustrated in table 5.

¹H-NMR (DMSO-d₆) δ ppm: 1.43 (3H, t, J=7.1Hz), 4.26-4.36 (2H, m), 7.05-7.15 (1H, m), 7.35-7.50 (3H, m), 7.90-8.25 (4H, m), 10.89 (1H, s)

25

Example 10

5,7-Difluoro-3-(5-piperazin-1-ylmethylfuran-2-carbonyl)-

aminobenzofuran-2-carboxamide (compound 9-1)

A solution of tert-butyl 4-[5-(2-carbamoyl-5,7-difluorobenzofuran-3-ylcarbamoyl)furan-2-ylmethyl]piperazine-1-carboxylate (compound 2-142, 0.44g) in 4mol/L hydrogen chloride in ethyl acetate (18mL) was stirred at room temperature for 4 hours. The solvent of the reaction mixture was removed under reduced pressure. The residue was dissolved in ethyl acetate, methanol and a 1mol/L aqueous solution of sodium hydroxide, and the mixture was stirred at room temperature for 20 minutes. Water and ethyl acetate were added to the mixture. The organic layer was separated, and the solvent was removed under reduced pressure to give the title compound (0.166g). The structural formula was illustrated in the table 5.

¹H-NMR (DMSO-d₆) δ ppm: 2.30-2.50 (4H, m), 2.65-2.80 (4H, m), 3.59 (2H, s), 6.50-6.65 (1H, m), 7.25-7.40 (1H, m), 7.50-7.65 (1H, m), 7.85-8.00 (1H, m), 8.06 (1H, brs), 8.39 (1H, brs)

Example 11

6-Chloro-3-(4-fluoro-3-morpholin-4-ylmethylbenzoylamino)-benzofuran-2-carboxamide (compound 10-1)

A mixture of 4-fluoro-3-methylbenzoic acid (0.1g), N-bromosuccinimide (0.121g), benzoyl peroxide (0.021g, 25% wet) and carbon tetrachloride was heated under reflux for 4 hours. Insoluble materials were removed by filtration. The filtrate was concentrated under reduced pressure, and the residue was purified by column chromatography on silica gel (eluent: hexane: ethyl acetate = 9:1-4:1:1% acetic acid was added) to give

3-bromomethyl-4-fluorobenzoic acid (0.151g).

N,N-Dimethylformamide (1 drop) was added to a mixture of 3-bromomethyl-4-fluorobenzoic acid and thionyl chloride (2mL) at room temperature, and the mixture was stirred at 80°C for 2.5 hours. The reaction mixture was concentrated under reduced pressure. The residue was dissolved in tetrahydrofuran (1mL), and the resulting solution was added to a mixture of 3-amino-6-chlorobenzofuran-2-carboxamide (0.1g), dimethylaniline (0.19mL) and tetrahydrofuran (2mL). The mixture was stirred at room temperature for 2 hours. Water was added to the reaction mixture, and the resulting precipitate was collected by filtration to give 3-(3-bromomethyl-4-fluorobenzoylamino)-6-chlorobenzofuran-2-carboxamide (0.042g).

A mixture of 3-(3-bromomethyl-4-fluorobenzoylamino)-6-chlorobenzofuran-2-carboxamide, morpholine (0.0092g), PS-NMM (registered mark) resin (argonaut, 100-200mesh, 1% divinylbenzene, 1.98mmol/g resin, 0.152g) and 1-methyl-2-pyrrolidone (2mL) was stirred at 80°C for 1 hour. After cooling the reaction mixture to room temperature, TMI-isocyanate (registered mark) (mimotope, D-type, 100micromol/unit, 2units) was added and the mixture was stirred at 80°C for 2 hours. The reaction mixture was purified by a SCX ion-exchanged column conditioned with methanol (argonaut, 2g, washing solvent: N,N-dimethylformamide, eluent: 2mol/L ammonia in methanol) to give the title compound (0.024g). The structural formula was illustrated in the table 5.

¹H-NMR (DMSO-d₆) δ ppm: 2.35-2.50 (4H, m), 3.50-3.70 (6H, m),

7.35-7.50 (2H, m), 7.70-7.80 (1H, m), 7.90-8.15 (3H, m),
8.15-8.35 (2H, m), 11.05 (1H, s)

MS (ESI, m/z) : 432 (M+H)+

5 Example 12

3-[5-(3-Dimethylaminomethylphenyl)furan-2-carbonyl]amino-
5,7-difluorobenzofuran-2-carboxamide (Compound 11-1)

A mixture of 3-[5-(3-chloromethylphenyl)furan-2-
carbonyl]amino-5,7-difluorobenzofuran-2-carboxamide

10 (compound 1-316, 0.041g), dimethylamine hydrochloride
(0.0282g), PS-NMM (registered mark) resin (argonauts,
100-200mesh, 1% divinylbenzene, 1.98mmol/g resin, 0.144g) and
1-methyl-2-pyrrolidone (2mL) was stirred at 80°C for 7.5 hours.
After cooling the reaction mixture to room temperature,
15 TMI-isocyanate (registered mark) (mimotope, D-type,
100micromol/unit, 3units) was added, the mixture was stirred
at 80°C for 4 hours. The mixture was purified by a SCX ion-exchanged
column conditioned with methanol (argonaut, 2g, washing
solvent: N,N-dimethylformamide, eluent: 2mol/L ammonia in
20 methanol) to give the title compound (0.0045g). The structural
formula was illustrated in table 5.

¹H-NMR (DMSO-d₆) δ ppm: 2.18 (6H, s), 3.47 (2H, s), 7.20-7.40
(2H, m), 7.40-7.55 (2H, m), 7.55-7.65 (1H, m), 7.75-7.90 (2H,
m), 7.95-8.10 (1H, m), 8.16 (1H, brs), 8.39 (1H, brs), 11.30
25 (1H, s)

MS (ESI, m/z) : 440 (M+H)+

N-(2-carbamoyl-5,7-difluorobenzofuran-3-yl)-6-(3-dimethyl
aminomethylphenoxy)nicotinamide (compound 11-2) was prepared
in a manner similar to those as described in example 12 using
N-(2-carbamoyl-5,7-difluorobenzofuran-3-yl)-6-

5 (3-chloromethylphenoxy)nicotinamide (compounds 1-357) instead
of 3-[5-(3-chloromethylphenyl)furan-2-carbonyl]-
amino-5,7-difluorobenzofuran-2-carboxamide (compound 1-316).
The structural formula was illustrated in table 5.

¹H-NMR (DMSO-d₆) δ ppm: 2.17 (6H, brs), 3.44 (2H, brs), 7.05-7.30
10 (4H, m), 7.35-7.80 (3H, m), 8.07 (1H, brs), 8.34 (1H, brs),
8.35-8.50 (1H, m), 8.75-8.85 (1H, m), 10.88 (1H, brs)

Table 5

Compound No.	Structure	Compound No.	Structure
4-1		8-1	
4-2		9-1	
5-1		10-1	
6-1		11-1	
6-2		11-2	
7-1			

Test Example 1

Measurement of antagonistic action for human adenosine A_{2A}

5 receptor

1) Preparation of human adenosine A_{2A} receptor-expressed cells

When Chinese hamster ovary-K1 (CHO-K1) cells were grown to subconfluence, they were treated with trypsin and suspended

in D-MEM/F-12 medium (Invitrogen) including 10% fetal bovine serum (Sankojunyaku). The expression vector of human adenosine A_{2A} receptor prepared in OPTI-MEM I (Invitrogen) and the transfection reagent of Lipofectamine 2000 (Invitrogen) were added to the cell suspension, and those cells were inoculated in poly-D-lysine-coated 96-well plates (5×10^4 cells/well). After incubating the plates for 24-30 hours at 37°C in a humidified atmosphere with 5% CO₂, they were used for the subsequent assays.

2) Measurement of the inhibitory effect on human adenosine A_{2A} receptor activity

A test compound was dissolved in dimethyl sulfoxide (DMSO) and the resulting solution was added to D-MEM/F-12 medium supplemented with 2.5 unit/mL adenosine deaminase (Carbiochem) and 30 μ M rolipram (Sigma), a phosphodiesterase inhibitor to prepare a solution of the test compound. Adenosine deaminase was used for the inactivation of adenosine derived from the cells.

The human adenosine A_{2A} -expressed cells inoculated in the 96-well plates were washed twice with 150 μ L/well of D-MEM/F-12 medium and a 50- μ L aliquot of the test compound solution was then added to each well of the plates. After incubating the cells for 10 minutes at 37°C in a humidified atmosphere with 5% CO₂, a 50- μ L aliquot of 1.2 nM 5'-N-ethylcarboxamide adenosine (NECA; Sigma) including 2.5 unit/mL adenosine deaminase and 30 μ M rolipram was added to each well of the plates and incubated for 25 minutes at 37°C in a humidified atmosphere with 5% CO₂. Stopping the reaction and the subsequent measurement of cAMP were

performed by use of a cAMP enzyme-linked immunosorbent assay kit (Applied Biosystem). The chemoluminescence was measured by a Microplate Luminometer TR717 (Applied Biosystem). The experiments were carried out in duplicate wells.

5 Inhibitory rate of a test compound for NECA-induced cAMP production was found by the following formula:

Inhibitory rate (%) = $[1 - \{ (\text{cAMP level with a test compound and NECA}) - (\text{cAMP level without a test compound and NECA}) \} / \{ (\text{cAMP level with NECA}) - (\text{cAMP level without a test compound and NECA}) \}]$

10 x 100

 Using a Km value for NECA found by the measurement in the same plate as that of test compounds, an inhibitory constant Ki was found by the following formula:

$$K_i = IC_{50} / \{1 + (6 \times 10^{-10} / K_m)\}$$

15 The results were shown in Table 6.

Table 6

Compound NO.	Ki value (nM)	Compound NO.	Ki value (nM)
1-1	8.5	1-150	2.2
1-2	4.1	1-153	9.2
1-15	6.5	1-155	8.1
1-16	0.9	1-156	9.8
1-36	6.4	1-159	3.3
1-91	6.5	1-161	7.0
1-108	9.3	1-213	2.1
1-109	2.7	1-249	4.6
1-119	1.9	1-275	0.5
1-121	0.8	2-1	8.0
1-125	1.2	2-7	8.6
1-133	3.2	2-129	1.6
1-138	4.5	2-138	2.0
1-141	9.3	2-142	2.0
1-145	2.7	6-2	1.1

Test example 2

Haloperidol-induced catalepsy in rats

5 The effect of a compound on drug-induced catalepsy in rats was evaluated. Parkinson's disease is a movement disorder reflecting a decline in striatal dopamine due to the degeneration of dopaminergic neuron arising from the substantia nigra. When haloperidol, a blocker of dopamine D2 receptor is administered

10 in rats, the drug induces catalepsy which is considered as one of the Parkinson's disease-like symptoms since haloperidol

blocks the neurotransmission of dopaminergic neuron.

Male rats (Crj: CD(SD)IGS, body weight: 210-260g, Charles River Japan, Inc.) were used for the experiments. Three to five rats for each group were tested for catalepsy 5 hours after

5 intraperitoneal administration of 1mg/kg haloperidol (Serenace (registered mark); Dainippon Pharma), a dopamine D2 antagonist. Catalepsy was measured using the horizontal bar test (Morelli and Chiara, "Eur. J. Pharmacol.", vol.117, p.179-185 (1985)). A compound (10 mg/kg) was dissolved in PEG400 and administered
10 orally 2 hours prior to the measurement of catalepsy. PEG400 was also administered orally to the rats for the control group. The front feet of the rats were placed on a horizontal bar (3 mm in diameter) suspended 10-12cm above a platform and their rear feet were placed on the platform. The degree of catalepsy
15 produced in the rats was measured by how long it took for their front feet to remove themselves from the bar. A maximum of 180 seconds was allowed. The results were shown in Table 7.

Table 7

Group	Duration of catalepsy (sec)
Control	180.0
Compound 1-2	91.6
Compound 1-16	116.4
Compound 1-121	48.2
Compound 1-155	38.3

20 The compounds of the present invention exhibited superior catalepsy improvement action, and are accordingly useful as a

medicament for treating a Parkinson's disease.

INDUSTRIAL APPLICABILITY

Compounds represented by general formula (I) of the present
5 invention exhibit potent adenosine A_{2A} receptor antagonistic
activities, and are accordingly useful for treating or preventing
diseases mediated by adenosine A_{2A} receptors such as motor
function disorders, depression, an anxiety disorder, cognitive
function disorders, cerebral ischemia disorder, restless legs
10 syndrome and the like.